

Clinical Trial Protocol

Iranian Registry of Clinical Trials

12 Sep 2026

Investigating the effect of rotator cuff trigger points dry needling on the stability and function of the upper limb in people with shoulder pain

Protocol summary

Study aim

The aim of this study was to investigate the effect of rotator cuff dry needling on the stability and function of the upper limb in people with shoulder pain.

Design

A clinical trial with controlled group, single blind, randomized

Settings and conduct

In this clinical trial study, 40 patients with shoulder pain who referred to clinics affiliated to Shiraz School of Rehabilitation Sciences will be divided into two groups using a random method: the first group of dry needling and The second group receive sham dry needling.

Participants/Inclusion and exclusion criteria

This study are perform in people with shoulder pain who have had at least 3 months of pain and they have at least one active trigger point in two the muscle is made up of four rotator cuff muscles. And if they have special injury in the shoulder and neck joints or prohibited from using dry needling, they will be excluded from the study.

Intervention groups

Dry needling Group: People with shoulder pain who receive three sessions of dry needling in one week of treatment (every other day), in the trigger points of the rotator cuff muscles (supraspinatus, infraspinatus, subscapularis and teres minor). Control group: people with shoulder pain who receive the sham dry needling in the said muscles, in three sessions; They receive the same as the treatment group. Samples recruited from rehabilitation clinics of Shiraz University of Medical Sciences.

Main outcome variables

Shoulder stability and upper extremity function

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20190909044734N4**

Registration date: **2023-08-20, 1402/05/29**

Registration timing: **prospective**

Last update: **2023-08-20, 1402/05/29**

Update count: **0**

Registration date

2023-08-20, 1402/05/29

Registrant information

Name

Leila Abbasi

Name of organization / entity

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2023-09-06, 1402/06/15

Expected recruitment end date

2023-10-07, 1402/07/15

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Investigating the effect of rotator cuff trigger points dry needling on the stability and function of the upper limb in people with shoulder pain

Public title

Investigating the effect of dry needling of the rotator cuff muscles on the stability and function of the upper limb in people with shoulder pain.	
Purpose	
Treatment	
Inclusion/Exclusion criteria	
Inclusion criteria:	
Presence of shoulder pain for at least three months	
Existence of trigger points with three characteristics: 1. Existence of a palpable knot in the skeletal muscle band.2. Existence of excessive points Irritable in taut band 3. Patients who report local or referred pain after mechanical stimulation of trigger points. (with the difference that latent trigger points become painful only after mechanical stimulation and active points even without stimulation. It is a mechanism that causes myofascial pain)	
The presence of at least one trigger point in two of the four rotator cuff muscles	
Exclusion criteria:	
History of shoulder and neck surgery	
Instability of the glenohumeral joint and serious diseases related to the joint of cervical vertebrae, ang glenohumeral joint	
History of corticosteroid injections or local anesthetics in the past year	
Symptoms indicating systemic disorders	
History of any physiotherapy in the last 3 months	
History of diagnosis or medical reports about the presence of myopathy and neuropathy	
Cognitive problems	
Patients with any limitations in dry needling treatment (fear of needles, patient dissatisfaction, lymphoedema, Acute and emergency conditions, patients with coagulation problems, immune system disorders, vascular problems, diabetes, epilepsy, allergy to metals, patients with mental disorders, use of anticoagulants)	
Use of anti-platelet, anti-coagulant, anti-pain and anti-inflammatory drugs in the last three months	
Cardiovascular, rheumatoid, and neurological problems	
History of rotator cuff tears	
Age	
From 18 years old to 45 years old	
Gender	
Both	
Phase	
N/A	
Groups that have been masked	
- Outcome assessor - Data analyser	
Sample size	
Target sample size: 40	
Randomization (investigator's opinion)	
Randomized	
Randomization description	
Using of randomization software, 40 numbers are considered for each case and the software defines 10 blocks of 4, from 4 to 2 people in the dry needling group and 2 people in sham dry needling. They are dried and thus 40 people are completed.	
Blinding (investigator's opinion)	
Single blinded	
Blinding description	
The person who will evaluate the patients and analyze	
	the data is completely unaware of the details of the patient grouping and the treatments performed.
Placebo	
Used	
Assignment	
Parallel	
Other design features	
Secondary Ids	
empty	
Ethics committees	
1	
Ethics committee	
Name of ethics committee	Ethics Committee in Research, Shiraz Faculty of Rehabilitation Sciences
Street address	Abiverdi Street 1, Chamran Blvd
City	Shiraz
Province	Fars
Postal code	1733-71345
Approval date	2023-07-26, 1402/05/04
Ethics committee reference number	IR.SUMS.REHAB.REC.1402.009
Health conditions studied	
1	
Description of health condition studied	Shoulder pain
ICD-10 code	M25.51
ICD-10 code description	Pain in shoulder
Primary outcomes	
1	
Description	Stability of shoulder
Timepoint	Before the intervention, after the end of the treatment and one week after the end of the treatment as a follow-up
Method of measurement	Closed Kinetic Chain Upper Extremity Stability Test
2	
Description	Upper extremity function
Timepoint	

Before the intervention, after the end of the treatment and one week after the end of the treatment as a follow-up

Method of measurement

Quick Disabilities of Arm, Shoulder & Hand questionnaire

3

Description

Grip strenght

Timepoint

Before the intervention, after the end of the treatment and one week after the end of the treatment as a follow-up

Method of measurement

Dynamometer device

4

Description

Shoulder pain

Timepoint

Before the intervention, after the end of the treatment and one week after the end of the treatment as a follow-up

Method of measurement

Visual Analogue Scale

5

Description

The strength of internal and external rotators of the shoulder joint

Timepoint

Before the intervention, after the end of the treatment and one week after the end of the treatment as a follow-up

Method of measurement

Dynamometer device

Secondary outcomes

empty

Intervention groups

1

Description

Intervention group: dry needling: people with shoulder pain who received 3 dry needling sessions per week according to the routine of the control group 3 times during a week of treatment (one day in between), in the trigger points of the rotator cuff muscles (supraspinatus, infraspinatus, subscapularis and teres minor).

Category

Rehabilitation

2

Description

Control group: sham dry needling group: people with shoulder pain who received dry needling in the rotator

cuff muscles (supraspinatus, infraspinatus, subscapularis and teres minor) and practically the needle does not enter the skin.

Category

Rehabilitation

Recruitment centers

1

Recruitment center

Name of recruitment center

Shiraz School of Rehabilitation

Full name of responsible person

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

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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

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Person responsible for general inquiries

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Sharing plan

Deidentified Individual Participant Data Set (IPD)

No - There is not a plan to make this available

Justification/reason for indecision/not sharing IPD

No more information

Study Protocol

No - There is not a plan to make this available

Statistical Analysis Plan

No - There is not a plan to make this available

Informed Consent Form

No - There is not a plan to make this available

Clinical Study Report

No - There is not a plan to make this available

Analytic Code

No - There is not a plan to make this available

Data Dictionary

No - There is not a plan to make this available