

Clinical Trial Protocol

Iranian Registry of Clinical Trials

11 Sep 2026

Effects of Sustained Natural Apophyseal Glides with and without Thoracic Postural Correction Techniques on Pain, Range of Motion and Disability in Patients with Mechanical Neck Pain

Protocol summary

Study aim

To determine the effects of Sustained Natural Apophyseal Glides with and without thoracic postural correction techniques on mechanical neck pain

Design

The study design will be a Randomized Controlled Trial with a parallel group design of 28 patients.

Settings and conduct

The study setting will be International Therapy Services ITS Clinic, Lahore. The sample would be taken by using a non-probability convenience sampling technique.

Participants/Inclusion and exclusion criteria

Inclusion Criteria: • Individuals having localized pain or stiffness in the spine or both combined b/w C3 to C7 without upper limbs radiculopathy • Patients with chronic neck pain >3 months • Pain reported on NPRS score ≥ 3 in the neck region with a limited or painful range
Exclusion Criteria: • Tuberculosis, carcinoma, heart disease, osteoporosis. • Neural disorders • Any trauma or localized infection in the neck region, cervical stenosis, metabolic diseases in bone and joint. • Hyper flexibility • Open sores • Ongoing radiotherapy, chemotherapy, steroid therapy, or anticoagulants. • Allergy to hot pack • Patients with a history of surgery in the cervical spine region within a year

Intervention groups

Group A will receive combination of a sustained facet glide with movement applied at the facet joints between cervical C3 to C7. Group B will receive sustained natural apophyseal glides same as group A along thoracic postural correction technique. Thoracic postural correction technique includes active as well as therapist-facilitated stretches.

Main outcome variables

Pain, Range of Motion and Disability

General information

Reason for update

Trial has been completed.

Acronym

IRCT registration information

IRCT registration number: **IRCT20190717044238N5**

Registration date: **2023-03-06, 1401/12/15**

Registration timing: **registered_while_recruiting**

Last update: **2023-09-24, 1402/07/02**

Update count: **1**

Registration date

2023-03-06, 1401/12/15

Registrant information

Name

Fareeha Amjad

Name of organization / entity

The University of Lahore

Country

Pakistan

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+92 42 99200600

Email address

fari_fairy22@yahoo.com

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2023-03-01, 1401/12/10

Expected recruitment end date

2023-06-30, 1402/04/09

Actual recruitment start date

2023-03-05, 1401/12/14

Actual recruitment end date

2023-06-25, 1402/04/04

Trial completion date

2023-08-05, 1402/05/14

Scientific title

Effects of Sustained Natural Apophyseal Glides with and without Thoracic Postural Correction Techniques on Pain, Range of Motion and Disability in Patients with Mechanical Neck Pain

Public title

Sustained Natural Apophyseal Glides and Thoracic Posture Correction Technique on Mechanical Neck Pain

Purpose

Treatment

Inclusion/Exclusion criteria**Inclusion criteria:**

Individuals having localized pain or stiffness in spine or both combined b/w C3 to C7 without upper limbs radiculopathy Patients with chronic neck pain >3 months Pain reported on NPRS score ≥3 in neck region with limited or painful range of motion Painful and limited cervical range of motion

Exclusion criteria:

Tuberculosis, carcinoma, heart disease, osteoporosis Neural disorders Any trauma or localized infection in neck region, cervical stenosis, metabolic diseases in bone and joint Hyper flexibility Open sores Ongoing radiotherapy, chemotherapy, steroid therapy or anticoagulants Allergy to hot pack Patients with history of surgery in cervical spine region with in a year

Age

From **18 years** old to **70 years** old

Gender

Both

Phase

3

Groups that have been masked

- Outcome assessor

Sample size

Target sample size: **28**

Actual sample size reached: **28**

Randomization (investigator's opinion)

Randomized

Randomization description

Patients will be assigned to either control or treatment group by random selection using a computer-generated table of random numbers. These numbers will be placed inside sealed envelopes. Another individual, who is not involved in the research, will then open these envelopes and assign the patients to their respective groups based on the instructions provided.

Blinding (investigator's opinion)

Single blinded

Blinding description

A skilled and experienced physiotherapist, who is not connected to the study, will making assessments on outcomes or gathering data on outcome variables.

Placebo

Not used

Assignment

Parallel

Other design features**Secondary Ids**

empty

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

Riphah College of Rehabilitation and Allied Health Sciences Lahore

Street address

F83G+V25, Madar-e-Millat Road, Quaid-e-Azam Industrial Estate Quaid e Azam Industrial Estate, Lahore, Punjab

City

Lahore

Postal code

54000

Approval date

2023-01-02, 1401/10/12

Ethics committee reference number

REC/RCR & AHS/23/0113

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

Mechanical neck pain

ICD-10 code

M54.2

ICD-10 code description

Cervicalgia

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

Functional disability

Timepoint

6 weeks

Method of measurement

Neck Disability Index is a self-report questionnaire designed to determine how neck pain affects a patient's daily life and the disability of patients with neck pain. It consists of 10 questions that ask about activities of daily living. More the score, the more significant the disability. The sample questionnaire is attached at the end. Four sections relate to subjective symptoms, and the remaining 6 sections relate to activities of daily living. Each unit is scored from 0 to 5 points, giving a maximum score of 50. The total score of the neck disability index ranges from 0 to 50 points

2**Description**

Pain
Timepoint
6 weeks
Method of measurement
Patient level of pain will be assessed using this scale.
This scale ranges from 0 to 10. 0 indicates "no pain" and 10 indicates "worst pain"

3

Description
Range of motion
Timepoint
6 weeks
Method of measurement
Changes from the Baseline ROM range of Motion of Cervical spine will be taken with the Help of Goniometer

Secondary outcomes

empty

Intervention groups

1

Description
Intervention group: Group A will receive conventional therapy (Hot pack and Transcutaneous Electrical Nerve Stimulation) along sustained natural apophyseal glides. Sustained facet glide with movement will be applied at the facet joint between cervical C3 to C7. It will be performed in sitting or standing position of the patient. Sustained Natural Apophyseal Glides involves application of accessory passive glide to cervical vertebrae by a physiotherapist while the patient will simultaneously perform an active movement. Glide given is in the direction of the plane of facet joints, and technique is usually performed in weight-bearing position like standing, sitting etc. Patients in control group will receive treatment for alternate days, 3 days a week for 6 weeks

Category
Treatment - Other

2

Description
Intervention group: Group B will receive conventional therapy same as group A with sustained natural apophyseal glides and thoracic posture correction technique. Thoracic postural correction technique includes active as well as therapist-facilitated stretches. Active stretches will be thoracic extension in sitting, Wall angle stretch and Corner stretch, while the therapist-facilitated stretches will be seated mid-thoracic stretch and prone mid thoracic stretch. Stretches will be maintained for 15- 20 seconds with 10 repetitions of each stretch per session. Patients in control group will receive treatment for alternate days, 3 days a week for 6 weeks.

Category
Treatment - Other

Recruitment centers

1

Recruitment center
Name of recruitment center
International Therapy Services
Full name of responsible person
Dr. Tayyab Malik
Street address
International Therapy Services UET Lahore
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Lahore
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Phone
+92 306 2149648
Email
itsclinicpak@gmail.com

Sponsors / Funding sources

1

Sponsor
Name of organization / entity
Riphah International University Lahore
Full name of responsible person
Dr. Fareeha Amjad
Street address
Riphah Quaid e Azam Campus, 28-M Quaid-e-Azam industrial Estate, kot lakhpat, Lahore
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Phone
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Email
fari_fairy22@yahoo.com
Grant name
Grant code / Reference number
Is the source of funding the same sponsor organization/entity?
No
Title of funding source
Riphah International University Lahore
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
Academic

Person responsible for general inquiries

Contact

Name of organization / entity
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Full name of responsible person
Rameen Qayyum
Position
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Person responsible for updating data

Contact

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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

No - There is not 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

Not applicable

Data Dictionary

Yes - There is a plan to make this available

Title and more details about the data/document

Consent Form in its original format with no information about any participant study protocol- how the intervention was given to both groups

When the data will become available and for how long

Data would be available after the completion of the research at the end of 2023

To whom data/document is available

People working in an academic and clinical setting can have access to the above-mentioned information/documents

Under which criteria data/document could be used

Data can only be used for Research Purposes

From where data/document is obtainable

Data can be asked for at the following email address: rameenqayyum95@gmail.com

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

One can ask for data at the given email address and it would be provided after knowing the general implications of sharing that particular data.

Comments