

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

10 Jun 2026

### Effectiveness of Acupressure with Stretching and Acupressure Technique alone in Patients with Perceived Stress Related Sub-Acute Neck Pain

#### Protocol summary

##### Study aim

To evaluate the effects of neck acupressure with and without stretching on perceived stress related sub-acute neck pain and to evaluate the effects of neck acupressure on disability, perceived stress and range of motion of the neck.

##### Design

Single blinded, parallel assigned, multi-centred, randomised clinical trial, quantitative study, sample size 48 (2 groups each with 24 participants) with perceived stress related sub-acute neck pain assessed by pain and stress, Simple random sampling technique

##### Settings and conduct

This study was conducted at physiotherapy department of Chatha Hospital Jaranwala. The study was completed in 4 months that included collection and analysis of data. Participants were kept anonymous or conducted in single blind trial

##### Participants/Inclusion and exclusion criteria

Included participants: Middle aged male and female participants, moderate score perceived stress scale score (14 or greater), with neck pain history of >30 days, complain of neck pain > 30mm on VAS, non-radiating neck pain and not undergoing psychiatry treatment. Excluded Participants: with any diagnosed musculoskeletal disorder, Pregnant females, or participants suffering from cancer, undergone surgery in past 3 months, with neck pain history of chronic, HIV disorders, swelling, skin issues, high blood pressure, any wound or contagious disease at the acupressure points, any neurological disorder, any vascular disorders and any infectious disease were excluded from study.

##### Intervention groups

participants were enrolled into 2 equal treatment groups, Group A (acupressure with stretching group) and Group B (acupressure alone group). Both groups with neck isometrics as baseline treatment

##### Main outcome variables

Primary Outcome Measure The cervical pain was

assessed by VAS Secondary Outcome Measure The cervical disability was measured by using the Neck Disability index and the range of motion was measured by goniometer

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20210806052089N1**

Registration date: **2021-09-07, 1400/06/16**

Registration timing: **registered\_while\_recruiting**

Last update: **2021-09-07, 1400/06/16**

Update count: **0**

##### Registration date

2021-09-07, 1400/06/16

##### Registrant information

##### Name

Fatima Naseem Bhatti

##### Name of organization / entity

The university of Faisalabad

##### Country

Pakistan

##### Phone

+92 41 4312868

##### Email address

fatimabhatti1765@yahoo.com

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2021-06-11, 1400/03/21

##### Expected recruitment end date

2021-12-22, 1400/10/01

##### Actual recruitment start date

empty

**Actual recruitment end date**  
empty

**Trial completion date**  
empty

**Scientific title**  
Effectiveness of Acupressure with Stretching and Acupressure Technique alone in Patients with Perceived Stress Related Sub-Acute Neck Pain

**Public title**  
Effectiveness of acupressure in patients with perceived stress related sub-acute neck pain.

**Purpose**  
Health service research

**Inclusion/Exclusion criteria**  
**Inclusion criteria:**  
Middle aged male and female participants between minimum age of 30 and maximum of 50 years  
Effectiveness of Acupressure with Stretching and Acupressure Technique alone in Patients with Perceived Stress Related Sub-Acute Neck Pain moderate score  
perceived stress scale score (14 or greater) Willing to participate by signing the consent form With neck pain history of >30 days (sub-acute) Complain of neck pain > 30mm on VAS, (moderate) Non-radiating neck pain Not undergoing psychiatry treatment or psychotropic medication were included into study  
**Exclusion criteria:**  
Participants with any diagnosed musculoskeletal disorder (spondylolisthesis, spondylitis, Rheumatoid arthritis etc.) Pregnant females, Participants suffering from cancer, Undergone surgery in past 3 months With neck pain history of >90 days (chronic) HIV disorders, swelling, skin issues High blood pressure Any wound or contagious disease at the acupressure points Any neurological disorder (Parkinson's, dementia, stroke etc.) Any vascular disorders (phlebitis, vertebral or carotid artery disorders, atherosclerosis etc.) Any infectious disease (T.B, Hepatitis etc.) were excluded from study

**Age**  
From **30 years** old to **50 years** old

**Gender**  
Both

**Phase**  
3

**Groups that have been masked**

- Participant

**Sample size**  
Target sample size: **48**

**Randomization (investigator's opinion)**  
Randomized

**Randomization description**  
Simple random sampling technique (probability sampling) where each participant has equal chance of getting enrolled into the study, was used for the sampling and after which the participants were enrolled into 2 equal treatment groups, Group A (acupressure with stretching group) and Group B (acupressure alone group) by lottery method where the selection of

participants into groups depends upon the card drawn and the specific number assigned to participants.

**Blinding (investigator's opinion)**

Single blinded

**Blinding description**

Participants will be aware of the description of treatments being performed in both groups but will not be aware which treatment will be performed on them. The participants of both groups will not be aware of the study groups. All the participants will be masked in the study. This will be carried out by keeping the anonymous by study period.

**Placebo**

Not used

**Assignment**

Parallel

**Other design features**

**Secondary Ids**

empty

**Ethics committees**

1

**Ethics committee**

**Name of ethics committee**

Ethics and technical committee of university of Faisalabad

**Street address**

The university of Faisalabad, university town, Sargodha road ,Faisalabad

**City**

Faisalabad

**Postal code**

38000

**Approval date**

2021-05-21, 1400/02/31

**Ethics committee reference number**

TUF/DR/SA/MSPP/2021/213-230

2

**Ethics committee**

**Name of ethics committee**

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**Street address**

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## Health conditions studied

### 1

#### Description of health condition studied

neck pain

#### ICD-10 code

M54.2

#### ICD-10 code description

Cervicalgia

## Primary outcomes

### 1

#### Description

Visual Analogous Scale

#### Timepoint

Baseline, after 2weeks(6th session), after 4weeks(12th session)

#### Method of measurement

This involves pain range from no pain to max unbearable pain. Here, the VAS is considered as a horizontal line that measures 100mm in length. The patient marks the line at the point they think the severity of their perception of pain can represent.

## Secondary outcomes

### 1

#### Description

Neck Disability Index

#### Timepoint

Baseline and after 4th week (12th session)

#### Method of measurement

Neck disability index (NDI) is a questionnaire used to assess the disability caused by the neck pain in everyday life. It involves 10 questions related to the activities of daily livings, that gives 6 possible options and point ranging 0-5, the least one is marked gives 0 score and last option being marked gives the 5 points, total points are 50 and the scoring is done by, dividing the patient score by 50 and multiplying by 100 gives a score in percent. If certain question (e.g. x) is missed because the patient doesn't do it, then total score will be divided by (50-x) points. The interpretation of NDI is given as: 0-4points (0-8%) no disability, 5-14points (10 - 28%) mild disability, 15-24points (30-48%) moderate disability, 25-34points (50- 64%) severe disability, 35-50points (70-100%) complete disability

### 2

#### Description

Perceived stress scale

#### Timepoint

Baseline and after 4th week (12th session)

#### Method of measurement

The perceived stress scale (PSS) is most commonly used psychometric scale for the measurement of perception of stress. This is simple questionnaire with simple and

related to nature questions. It describes the feeling of patient regarding certain aspects in last month. The scores of PSS are obtained by simply reversing the patient's responses of positive sensed questions "4, 5, 7 and 8". i.e. (0=4, 1=3, 2=2, 3=2, 4=0) and then summing up all responses. The score can range from 0-40 where the higher scores represent higher level of stress by patient. The scoring is interpreted as: 0-13= low perceived stress, 14-26= moderate perceived stress and 27-40= high perceived stress.

### 3

#### Description

Cervical range of motion

#### Timepoint

Baseline and after 4th week (12th session)

#### Method of measurement

The cervical ranges of motion (flexion, extension, side bending and rotation) was measured by using Goniometer.

## Intervention groups

### 1

#### Description

Intervention group: Group A (acupressure with stretching). The group received warm-up session of hot-pack and neck isometric exercises with 10s hold at 20% of the isometric strength repeated for 8 repetitions in 4 all 4 directions. Participants in group A received acupressure at 21 acupressure points, Each of the acupoint was pressed for 10 seconds and repeated for 5 times. This acupressure treatment took almost 20 minutes of session and then group A participants received addition of stretching. Stretching techniques were performed in such order: towards lateral flexion (upper part of trapezius), ipsilateral flexion and rotation (scalene) and towards flexion (extensor muscles) each for 30 sec and repeated 2-3 times. This complete session took 30 minutes total.

#### Category

Treatment - Other

### 2

#### Description

Intervention group: Group B (Acupressure alone group). The participants of group B received warm-up of hot-pack and isometric exercise and then the acupressure therapy on all points for treatment time of 20 minutes only was performed.

#### Category

Treatment - Other

## Recruitment centers

### 1

#### Recruitment center

Name of recruitment center

Chatha Hospital Jaranwala  
**Full name of responsible person**  
Dr.Hareem Aslam  
**Street address**  
Main road ,Faisalabad road  
**City**  
Jaranwala  
**Postal code**  
38000  
**Phone**  
+92 41 4711058  
**Email**  
Zayina.pearl.zp@gmail.com

## Sponsors / Funding sources

### 1

#### Sponsor

**Name of organization / entity**  
Yasmin physiotherapy centre  
**Full name of responsible person**  
Osama Ramzan  
**Street address**  
Sargodha road, near muslim town3 opposite KIA  
motors  
**City**  
Faisalabad  
**Postal code**  
38000  
**Phone**  
+92 41 8785675  
**Email**  
Osamaramzan@gmail.com  
**Web page address**  
Http://omiphysio.com

#### Grant name

**Grant code / Reference number**

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

Yes

#### Title of funding source

Yasmin physiotherapy centre

#### 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**  
Fatima Naseem Bhatti

**Position**  
Student  
**Latest degree**  
Master  
**Other areas of specialty/work**  
Physiotherapy  
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## Person responsible for scientific inquiries

#### Contact

**Name of organization / entity**  
The university of Faisalabad  
**Full name of responsible person**  
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**Position**  
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## Person responsible for updating data

#### Contact

**Name of organization / entity**  
The university of Faisalabad  
**Full name of responsible person**  
Fatima Naseem Bhatti  
**Position**  
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**Latest degree**  
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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**

As we have signed and assured the participants that there data will not be shared anywhere else other than this study

**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**

Yes - There is a plan to make this available

**Analytic Code**

Not applicable

**Data Dictionary**

Not applicable

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

Effectiveness of Acupressure with Stretching and Acupressure Technique alone in Patients with Perceived Stress Related Sub-Acute Neck Pain Everything related to the trial would be provided study protocol, primary and secondary outcome measure data will be shared no further details regarding personal information of the patient will be shared.

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

Starting from January 2022

**To whom data/document is available**

Anyone relevant to the medical field can apply for it

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

Medical students , researchers and anyother healthcare related person can ask for it

**From where data/document is obtainable**

By Email

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

An Email stating the use of data will be appreciated

**Comments**