

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

10 Jun 2026

### Effectiveness of Low Level Laser Therapy for treatment of cervical pain in patients with trigger points of upper trapezius

#### Protocol summary

##### Study aim

To determine the effectiveness of low level laser therapy for treatment of cervical pain in patients with trigger point of upper trapezius muscle.

##### Design

Single Blind , parallel , Randomized Controlled Trial

##### Settings and conduct

This Randomized Controlled Trial ,Double Blind ,Parallel study will be conducted at University of Lahore Teaching Hospital

##### Participants/Inclusion and exclusion criteria

Inclusion : Patients diagnosed with active upper trapezius trigger point. •Both Male and female •Age ranges from 18 to 55 Exclusion : Participants having contraindications for LASER e.g. Pregnancy, Tumors, Implants, Pacemaker, •Past History of fracture of shoulder joint or cervical spine •Whiplash injury •Comorbid medical diagnosis e.g. osteoarthritis •Psychiatric illness (e.g., schizophrenia or substance abuse) •Irritable skin around the area of treatment

##### Intervention groups

Patients of group-A will receive conventional physical therapy along with low level laser therapy ( LLLT with Omega Class IIIB Laser therapy unit with pen probe.) While patients of group B, will receive conventional physical therapy only.

##### Main outcome variables

Neck pain , Neck Range of Motion and Neck Disability Indexes

#### General information

##### Reason for update

##### Acronym

Cervical pain Trigger points Upper trapezius Neck Pain Neck Range of Motion Neck Disability Index Low Level Laser Therapy

##### IRCT registration information

IRCT registration number: **IRCT20210409050912N1**

Registration date: **2021-08-29, 1400/06/07**

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

Last update: **2021-08-29, 1400/06/07**

Update count: **0**

##### Registration date

2021-08-29, 1400/06/07

##### Registrant information

###### Name

Iqra Waseem

###### Name of organization / entity

Faisal Institute of Health Sciences

###### Country

Pakistan

###### Phone

+92 41 8818334

###### Email address

iqra.waseem91@gmail.com

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2021-07-28, 1400/05/06

##### Expected recruitment end date

2021-12-25, 1400/10/04

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

##### Scientific title

Effectiveness of Low Level Laser Therapy for treatment of cervical pain in patients with trigger points of upper trapezius

##### Public title

Low Level Laser Therapy for treatment of cervical pain in patients with trigger points of upper trapezius

### **Purpose**

Treatment

### **Inclusion/Exclusion criteria**

#### **Inclusion criteria:**

Diagnosed patients with active upper trapezius trigger point Age range from 18-55 years Male and Females

#### **Exclusion criteria:**

Participants having contraindications for LASER e.g. Pregnancy, Tumors, Implants, Pacemaker, Irritable skin around the area of treatment Past History of fracture of shoulder joint or cervical spine Whiplash injury Comorbid medical diagnosis e.g. osteoarthritis Psychiatric illness (e.g., schizophrenia or substance abuse). Psychiatric illness (e.g., schizophrenia or substance abuse).

### **Age**

From **18 years** old to **55 years** old

### **Gender**

Both

### **Phase**

3

### **Groups that have been masked**

- Outcome assessor

### **Sample size**

Target sample size: **54**

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

Randomized

### **Randomization description**

Participants will be randomly allocated by use of computer-generated randomization sheet into two groups One for Group A and other for B. 2 Sets of 27 Unique Numbers per Set Range: From 1 to 54

### **Blinding (investigator's opinion)**

Single blinded

### **Blinding description**

The outcome assessor will be blind and will assess the subjects for inclusion criteria and will make assessment at the baseline ,then after 2 weeks and than after 4 weeks . Then outcome assessor will give all the recorded data to the principal investigator

### **Placebo**

Used

### **Assignment**

Parallel

### **Other design features**

## **Secondary Ids**

empty

## **Ethics committees**

### 1

#### **Ethics committee**

##### **Name of ethics committee**

The University of Lahore

##### **Street address**

Lahore

### **City**

Lahore

### **Postal code**

38000

### **Approval date**

2020-03-07, 1398/12/17

### **Ethics committee reference number**

IRB-UOL-FAHS/718-IX/2020

## **Health conditions studied**

### 1

#### **Description of health condition studied**

Upper Trapezius Trigger Points

#### **ICD-10 code**

Y93

#### **ICD-10 code description**

Trigger Points , Neck pain, neck ROM , Neck Functioning

## **Primary outcomes**

### 1

#### **Description**

Neck pain

#### **Timepoint**

It will be assessed three times ,First at Baseline , Second after 02 weeks of treatment and third and last at the end of 04 weeks of treatment .

#### **Method of measurement**

Numeric Pain Rating Score- The numerical pain rating scale is considered as one of the best method available for the estimation of the intensity of pain. The NPRS is a categorical scale which provides a continuous scale for magnitude estimation of pain. NPRS consists of 11 points, which range from 0-10.0=no pain, 1-4=mild pain, 4-7=moderate pain 7-10=severe pain.

## **Secondary outcomes**

### 1

#### **Description**

Neck Range of Motion.

#### **Timepoint**

First it will be assessed three times at Baseline , then after 02 weeks and at the end of 04 weeks of treatment .

#### **Method of measurement**

Universal Goniometer will used for measurement of Range of Motions of neck in all sides .

### 2

#### **Description**

Neck disability Index

#### **Timepoint**

First it will be assessed three times at Baseline , then after 02 weeks and at the end of 04 weeks of treatment.

#### **Method of measurement**

Neck Disability Index- The Neck Disability Index (NDI) is a 10-item questionnaire that measures a patient's self-





For sake of help and research promotion .

**From where data/document is obtainable**

from principal Investigator

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

The email is provided for this purpose .

**Comments**

This trial is purely based for study purpose .It has not involved any third party and any funding except educational support from the relative institute .