

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

10 Jun 2026

### “Comparison of extracorporeal shockwave and high intensity laser in reducing pain and improving function in chronic plantar fasciitis.”

#### Protocol summary

##### Study aim

By knowing the short and long term effects of ESWT(Extracorporeal shockwave therapy) and HILT(High intensity laser therapy) in patients of chronic plantar fasciitis, we can avoid the need of cortisone injection and surgery.

##### Design

A randomized controlled not blind trial. Samples will be divided into 2 groups.1 group will be treated with extra corporeal shock wave and 2nd will be treated with high intensity LASER therapy. Both groups will receive treatment for 3 weeks and than follow up will be taken after 2 months.

##### Settings and conduct

Setting: Lifeline Healthcare and Pain center, Opposite Jinnah hospital Lahore, Pakistan.

##### Participants/Inclusion and exclusion criteria

Inclusion criteria: both genders, minimum age 18 and maximum 70, VAS >5 on first step in morning, Informed consent, Symptoms duration more than 6 months.

Exclusion criteria: Previous surgery of plantar fasciitis, Pregnancy, Pacemaker fitted, Bleeding disorder, Any tumor of treatment site, calcaneous stress fracture checked by squeeze test.

##### Intervention groups

Patients will be randomly divided into two groups 15 subjects in each group. After taking baseline assessment Group 1 will be treated with BTL-6000 Shockwave therapy Intensity= 2.0 Bar, Frequency =10Hz, Number of Shocks=2000. The ESWT will be applied in a circular motion on the insertion site of the plantar fascia (1,000 shocks) and along the fascia (1,000 shocks). (9) twice a week for 3 weeks. Group 2 will receive Diowave 60W Class IV Laser three times per week for 3 weeks for a total of 9 sessions at wavelength of 980nm, a laser output of 30 Watts and energy dose of 10,000 Joules. Scanning method will be used. Both groups will be given some exercises to follow for home plan.

##### Main outcome variables

Pain (VAS) Function of foot by Foot functional Index

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20200317046798N1**

Registration date: **2020-04-17, 1399/01/29**

Registration timing: **prospective**

Last update: **2020-04-17, 1399/01/29**

Update count: **0**

##### Registration date

2020-04-17, 1399/01/29

##### Registrant information

##### Name

Saba Riaz

##### Name of organization / entity

Riphah International University Lahore, Pakistan

##### Country

Pakistan

##### Phone

+92 42 35225100

##### Email address

sabariaz317@gmail.com

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2020-05-10, 1399/02/21

##### Expected recruitment end date

2020-06-30, 1399/04/10

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

**Trial completion date**

empty

**Scientific title**

“Comparison of extracorporeal shockwave and high intensity laser in reducing pain and improving function in chronic plantar fasciitis.”

**Public title**

“Comparison of extracorporeal shockwave and high intensity laser in reducing pain and improving function in chronic plantar fasciitis.”

**Purpose**

Treatment

**Inclusion/Exclusion criteria****Inclusion criteria:**

Both genders minimum age 18 and maximum 70 Visual analogue Scale >5 on first step in morning Informed consent Symptoms duration more than 6 months

**Exclusion criteria:**

Previous surgery of plantar fasciitis, Pregnancy Pacemaker fitted Bleeding disorder, Any tumor of treatment site calcaneous stress fracture checked by squeeze test

**Age**

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

**Gender**

Both

**Phase**

4

**Groups that have been masked**

*No information*

**Sample size**

Target sample size: **30**

**Randomization (investigator's opinion)**

Randomized

**Randomization description**

Randomization by lottery method. Allocation concealment through sealed envelopes

**Blinding (investigator's opinion)**

Not blinded

**Blinding description****Placebo**

Not used

**Assignment**

Parallel

**Other design features**

Non-probability convenient sampling technique will be used to recruit the individuals for the study and then randomization will be done by lottery method to divide the individuals into 2 treatment groups. Sealed envelopes will be used for allocation concealment.

**Secondary Ids**

empty

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

Riphah College of Rehabilitation Sciences (RCRS)  
Research Ethical Committee (REC)

**Street address**

near Hajj Complex, I-14, Islamabad, Islamabad Capital Territory

**City**

Islamabaad

**Postal code**

46000

**Approval date**

2019-10-09, 1398/07/17

**Ethics committee reference number**

RCRS-RE-MS-OMPT/SPring 19/ 033

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

Plantar Fasciitis is one of the most common causes of heel pain. It involves inflammation of a thick band of tissue that runs across the bottom of your foot and connects your heel bone to your toes

**ICD-10 code**

M72.2

**ICD-10 code description**

Plantar fascial fibromatosis

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

Pain

**Timepoint**

before treatment. than after treatment (3 weeks), Follow Up (after 2 months)

**Method of measurement**

Visual Analogue scale , Foot functional Index (Short version)

**2****Description**

Functional Status

**Timepoint**

before treatment. than after treatment (3 weeks), Follow Up (after 2 months)

**Method of measurement**

Foot functional Index (Short version)

**Secondary outcomes**

empty

**Intervention groups****1****Description**

After taking baseline assessment Group 1 will be treated

with BTL-6000 Extracorporeal Shockwave therapy Intensity= 2.0 Bar, Frequency =10 Hz , Number of Shocks=2000. The ESWT will be applied in a circular motion on the insertion site of the plantar fascia (1,000 shocks) and along the fascia (1,000 shocks). twice a week for 3 weeks

**Category**

Treatment - Devices

**2****Description**

After taking baseline assessment group 2 will receive Diowave 60W Class IV Laser three times per week for 3 weeks for a total of 9 sessions at wavelength of 980 nm , a laser output of 30 Watts and energy dose of 10,000 Joules. Scanning method will be used.

**Category**

Treatment - Devices

**Recruitment centers****1****Recruitment center****Name of recruitment center**

Lifeline Health Care and Pain Center , opposite Jinnah hospital Lahore, Pakistan

**Full name of responsible person**

Saba Riaz

**Street address**

150-HBFC, Opp. Emergency Gate Jinnah Hospital,Lahore

**City**

Lahore

**Postal code**

54770

**Phone**

+92 42 35236690

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sabariaz317@gmail.com

**Web page address**

<http://www.lifelinehealthcare.org/>

**Sponsors / Funding sources****1****Sponsor****Name of organization / entity**

Lifeline Healthcare and pain center

**Full name of responsible person**

Saba Riaz

**Street address**

150-HBFC, opp Jinnah Hospital Emergency Gate, Lahore Pakistan

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Lahore

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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**

Lifeline Healthcare and pain center

**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**

Riphah International University Lahore, Pakistan

**Full name of responsible person**

Saba Riaz

**Position**

Student

**Latest degree**

Bachelor

**Other areas of specialty/work**

Physiotherapy

**Street address**

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**Full name of responsible person**

SABA RIAZ

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student  
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## Person responsible for updating data

### Contact

**Name of organization / entity**  
Riphah College of Rehabilitation Sciences (RCRS)  
**Full name of responsible person**  
Saba Riaz  
**Position**  
student  
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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

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

Yes - There is a plan to make this available

### Data Dictionary

Yes - There is a plan to make this available

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

study protocol, Statistical analysis, results

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

after completion of study

### To whom data/document is available

People working in Healthcare providing centers and academic institutes regarding healthcare.

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

According to Profession

### From where data/document is obtainable

Different search sites

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

Registration on specific site

### Comments