

Clinical Trial Protocol

Iranian Registry of Clinical Trials

13 Jun 2026

Comparing the efficiency of diode laser 940 nm and Er, Cr:YSGG 2780 nm on clinical indices and MMP-13 level during phase I periodontal therapy of chronic periodontitis

Protocol summary

Study aim

the effect of 2 wave-length laser therapy with SRP on GCF MMP-13 levels in chronic periodontitis

Design

clinical trial , double-blind, split-mouth

Settings and conduct

periodontics department, Islamic Azad university of medical science split-mouth design on 16 patients with 6 months follow up double blind study(participant and investigator)

Participants/Inclusion and exclusion criteria

No systemic disease, having at least 16 teeth, no use of drugs (anti-biotic, corticostroid, immuo-suppressive), no history of periodontal treatment during last 6 months, no pregnancy and/or breast-feeding

Intervention groups

test groups: 2 quadrants will be treated by diode laser 940nm or Er,Cr:YSGG 2780nm following SRP(scaling and root planing). control group: the third quadrant will recieve only SRP treatment

Main outcome variables

Matrix metalloproteinase(MMP)13 level clinical attachment level (CAL)

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20140318017053N9**

Registration date: **2019-02-14, 1397/11/25**

Registration timing: **retrospective**

Last update: **2019-02-14, 1397/11/25**

Update count: **0**

Registration date

2019-02-14, 1397/11/25

Registrant information

Name

Ferena Sayar

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 21 2256 4571

Email address

f_sayar@dentaliau.ac.ir

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2018-09-23, 1397/07/01

Expected recruitment end date

2018-11-22, 1397/09/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparing the efficiency of diode laser 940 nm and Er, Cr:YSGG 2780 nm on clinical indices and MMP-13 level during phase I periodontal therapy of chronic periodontitis

Public title

The effect of laser therapy on MMP-13 level during periodontal treatment

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria:

systemic health Having 16 teeth or more

Exclusion criteria:

smoking pregnancy breast-feeding use of drugs (corticosteroid, anti-biotic, immuno-suppressive) periodontal therapy during the last 6 months.

Age

From **25 years** old

Gender

Both

Phase

N/A

Groups that have been masked

- Participant
- Investigator

Sample size

Target sample size: **16**

More than 1 sample in each individual

Number of samples in each individual: **3**
collecting GCF

Randomization (investigator's opinion)

Not randomized

Randomization description**Blinding (investigator's opinion)**

Double blinded

Blinding description

investigator will not aware of the control and test sites, she only records clinical data. because of application of 2 wave-length laser, participants will not aware of type of received treatment in each quadrant

Placebo

Not used

Assignment

Parallel

Other design features

split-mouth

Secondary Ids

empty

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

Ethics committee of Faculty of Dentistry Tehran
Medical sciences , Islamic Azad University

Street address

No.4 , 10th Neyestan st. , Pasdaran Ave.

City

Tehran

Province

Tehran

Postal code

19585/175

Approval date

2018-04-22, 1397/02/02

Ethics committee reference number

IR.IAU.DENTAL.REC.1397.020

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

chronic periodontitis

ICD-10 code

K05.3

ICD-10 code description

Chronic periodontitis

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

clinical attachment level.

Timepoint

base-line , 2 and 6 months after phase I periodontal therapy.

Method of measurement

periodontal probe.

2**Description**

matrix metalloproteinase-13

Timepoint

base-line , 2 and 6 months after phase I periodontal therapy.

Method of measurement

ELISA test

Secondary outcomes**1****Description**

pocket depth.

Timepoint

base-line , 2 and 6 months after phase I periodontal therapy.

Method of measurement

periodontal probe.

2**Description**

bleeding on probing.

Timepoint

base-line , 2 and 6 months after phase I periodontal therapy.

Method of measurement

periodontal probe.

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

Intervention group 1: diode laser 940 nm (Ezlase, biolase, USA) , 1 W , 20 sec.

Category

Treatment - Devices

2**Description**

Intervention group 2 : laser Er,Cr:YSGG 2780 nm (iPlus, Waterlase, Biolase, USA) 1.5 W , 30 Hz , 11% air , 20% water , pulse duration 140 µs.

Category

Treatment - Devices

3**Description**

Control group: phase I periodontal therapy

Category

Treatment - Devices

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

Department of Periodontics, Faculty of Dentistry,
Tehran Medical sciences, Islamic Azad University

Full name of responsible person

DR.Ferena Sayar

Street address

department of periodontics, no:4, 10th neyestan st.,
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Web page address**Sponsors / Funding sources****1****Sponsor****Name of organization / entity**

Islamic Azad University

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

Islamic Azad University

Proportion provided by this source

1

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**

Islamic Azad University

Full name of responsible person

Ferena Sayar

Position

Associate professor

Latest degree

Specialist

Other areas of specialty/work

Dentistry

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Person responsible for scientific inquiries**Contact****Name of organization / entity**

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Position

Associate professor

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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

There is no further 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