



EFFECT OF LOW-LEVEL LASER THERAPY ON PAIN EXPERIENCED DURING LEVELING AND ALIGNMENT OF LOWER ANTERIOR TEETH: A RANDOMIZED CONTROLLED CLINICAL STUDY

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ABSTRACT

Objective: This randomized controlled clinical study investigated the effect of a 635 nm Gallium-Aluminum-Arsenide (Ga-Al-As) laser on pain perception during the leveling and alignment of lower anterior teeth. **Subjects and Methods:** Twenty eight orthodontic patients, 17 females and 11 males, were randomly selected who underwent leveling and alignment through a non-extraction approach. They were randomly divided according to into two groups: non-laser group and low-level laser therapy (LLLT) group. Both groups utilized the same leveling 0.012 inch Ni-Ti arch wires. The laser group was exposed to Ga-Al-As semiconductor diode laser with 635 nm wavelength, 6.5J/cm² energy density, for 10 seconds on 10 points distributed over the labial and lingual aspects of each root of the lower anterior teeth. This was applied immediately after archwire insertion and then at days 3,7,14, 28 of the first month. Each patient was provided with a Visual Analog Scale (VAS) to record the pain score at 4 hours, 6 hours, 24 hours, 3 days, 7 days, and 28 days. **Results:** In both laser and non-laser groups, there is a statistically non-significant increase of pain measurements at 6 and 24 hours and a non-significant decrease in mean of pain measurements after day 3. **Conclusions:** The LLLT, with current parameters and protocol, has a negligible effect on pain experienced during initial orthodontic leveling and alignment stage.

Keywords: Low-Level Laser Therapy, Orthodontic Leveling and Alignment, Pain, Visual Analog Scale

INTRODUCTION

Orthodontic therapy is essential for functional and several aesthetic rehabilitation therapies of the stomatognathic system. It is known that application of orthodontic force to the dental system stimulates a transient inflammatory process mediated by inflammatory mediators, with no pathological condition producing some sort of pain⁽¹⁾.Doubtless orthodontic pain is one of the unwanted side effects given a great concern among orthodontic patients directing them to their withdrawal from the treatment.Orthodontic pain is described as discomfort, dull pain, and hypersensitivity in affected teeth⁽²⁾, it usually reaches its peak 24 hours after wire engagement, and begins to reduce 72 hours after wire engagement⁽³⁾.

Several methods have shown a reduction of the discomfort caused by pain during tooth movement such as anti-inflammatory medication⁽⁴⁾, low-level laser therapy (LLLT), however, the secondary effects of the administration of non-steroidalanti-inflammatory drugs (NSAID) may affect the rate of tooth movement.⁽⁴⁾

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Low-level laser therapy (LLLT) has an energy output that is low enough so as not to cause the temperature of the treated tissues to rise above 36.5°C or normal body temperature. The LLLT has been introduced into medicine and dentistry for various clinical practices for its bio-stimulatory effect which contributed to wound healing, muscle relaxation, nerve regeneration, collagen synthesis, fibroblast proliferation, acceleration of bone regeneration, reduction of inflammation, to improve blood circulation and increase cell activity. In the field of orthodontics, researchers proposed a positive effect of LLLT in managing orthodontic pain and increasing bone deposition in midpalatal suture during rapid maxillary expansion in animals⁽⁵⁻⁸⁾.

Because of easy application and noninvasive use of LLLT in addition to its limited side effects and fewer contraindications, many studies used it to control pain resulted from orthodontic tooth movement reporting that LLLT was able to control pain during orthodontic tooth movement, however, other studies indicate that laser cannot⁽³⁾. The debate continues, as conflicting conclusions were reached in recent systematic reviews regarding clinical trials of LLLT therapy with orthodontic tooth movement (OTM), it appears that the findings reports revealed controversial results the effect of LLLT on orthodontic pain⁽⁹⁾. The extensive variability in the experimental designs and laser specs e.g. wavelength, power output, energy density, mode, duration, and frequency of laser applications is a major contributor factor to the existing conflicting conclusions and presents a challenge that should be considered⁽⁹⁾.

Since few studies are available investigated multiple doses of LLLT through leveling and alignment phase of orthodontic treatment⁽¹⁰⁾, the aim of the current study was directed to evaluate the effect of continuous multiple doses of LLLT on orthodontic pain during leveling and alignment of the lower anterior teeth. The hypothesis of the study was supposed that LLLT could reduce pain during leveling and alignment of the lower anterior teeth.

SUBJECTS AND METHODS

This randomized controlled clinical trial was performed from January 2018 to July 2018 on a total sample of 28 patients, 17 females and 11 males were selected from outpatient clinic, Department of Orthodontics, Faculty of Dental Medicine (Boys), Al-Azhar University, Cairo, Egypt. Institutional Review Board and Ethical Committee of Al-Azhar University reviewed and approved the study protocol. All patients and/or their parents who agreed to participate in this research signed an informed consent document that authorized their data to be used for research purposes. Based on the previous studies⁽¹⁰⁻¹³⁾, a power analysis using G*Power software (version 3.1.9.2; Universitat Dusseldorf, Dusseldorf, Germany) estimated that the sample size of 28 patients ensured more than 80% power to detect significant differences at a 0.05 significance level. Theywere randomly divided into two groups according to the intervention into laser group and non-laser group (Fig.1).

All patients were selected according the following inclusion criteria: Class I malocclusion, with moderate mandibular crowding, orthodontic treatment with no extraction on the mandibular arch, all mandibular teeth erupted (third molars not included), and no spaces in the mandibular arch and the following exclusion criteria were considered: use of analgesics, treatment requires extractions of any tooth, impacted or unerupted permanent teeth, treatment with intraoral or extraoral removable appliances, patients with cleft lip and palate, anomalies, and syndromes.

Every participant was treated with the same Roth pre-adjusted metallic brackets (3M Unitek, Monrovia, Calif), leveling and alignment for both groups were carried out starting with 0.012-inch Ni-Ti archwire (Ortho Organizer[®] Super Elastic TitaniumArchwiress, USA) for the 1st month and ligated with elastomeric ligatures. Laser group received low-level laser therapy (LLLT) via Ga-Al-As diode laser (SmartTM PRO, Lasotronix, Poland) with a wavelength of 635 nm, energy density of 6.5J\cm², power output 20 mW (Fig.2a).



FIG (1) CONSORT Flow Diagram.



FIG (2a) Gallium Aluminum Arsenide semiconductor diode laser equipment used in the study.



FIG (2b) Points of application of LLLT from the labial aspect.

This was applied on 10 points on each root of the lower anterior teeth 5 labial and 5 lingual (Fig.2b) for 10 sec (2 points: mesial and distal to the apical area and 2 points: mesial and distal to the cervical area and one on the middle of the root). These parameters and protocol of application delivered a 0.2J per point, 2J per tooth and 12 Joules as a total dose of energy was delivered to the lower anterior segment per session. This was doneon days 0, 3, 7 and 14 starting from the day of the archwire placement.

All patients requested to mark pain intensity represented from 0 to 10 on the Visual Analogue Scale (VAS) at 4 hours, 6 hours , 24 hours, 3 days, 1 week, and 28 days⁽¹¹⁾ (Fig.3), using the terms no pain (0) and the highest pain (10) perceived over first month during incising and rest.





Statistical Analysis

All VAS scores were statistically analyzed using the SPSS (Statistical Package for Social Sciences (SPSS, version 22, Inc, Chicago, III) program. Mann Whitney test was used to compare pain scores between groups.

RESULTS

Although 2 patients (1 male and 1 female) were dropped out due to either missing appointments of laser application or repeated breakage of the orthodontic appliance the remaining 26 patients (16 females and 10 males) had completed the course of this study.

A comparison between the two studied groups (table 1) during incising shows a statistically nonsignificant increase at 6 and 24 hours and decrease in mean degree of pain measurements after day 3. A comparison between the two studied groups (table 2) during restshows a statistically nonsignificant increase at 6 and 24 hours and decrease in mean degree of pain measurements after day 3.

TABLE (1) Comparison between the two studied groups during incising

	Non-Laser (n = 13)		laser (n = 13)		T	n				
	Mean	±SD	Mean	±SD		P				
Incising										
4hr	4.50	0.71	3.30	1.42	46.0	0.569				
6hr	5.30	0.95	4.90	1.66	44.0	0.674				
24hr	5.20	1.32	4.80	1.23	43.0	0.543				
3Day	3.90	0.57	3.60	1.58	41.0	0.651				
7Day	2.60	1.29	2.10	2.27	44.0	0.684				
28 days	2.10	1.07	1.60	1.73	43.0	0.631				

U: Mann Whitney test

p: *p*-value for comparing between the studied groups *: Statistically significant at $p \le 0.05$

Table (2):Comparison between the twostudied groups during rest

	Non-Laser (n = 13)		laser (n = 13)		U	n				
	Mean	±SD	Mean	±SD	U	r				
Rest										
4hr	4.70	1.49	3.90	1.45	36.50	0.315				
6hr	5.0	1.56	3.80	1.40	28.50	0.105				
24hr	4.70	1.57	4.60	2.22	40.0	0.481				
3Day	2.10	1.66	1.70	1.0	32.50	0.195				
7Day	0. 90	0.26	0.60	0.20	33.50	0.145				
28 days	0.60	0.40	0.60	0.40	29.0	0.123				

U: Mann Whitney test

p: *p*-value for comparing between the studied groups *: Statistically significant at $p \le 0.05$

DISCUSSION

It is apparent that almost all patients submitted to fixed orthodontic treatment suffer some type of discomfort, considering the separation of teeth for posterior orthodontic banding or after the archwire insertion, which may also discourage them to continue the treatment, or even giving up at the beginning of the process⁽¹⁰⁾. The perception of pain varies considerably from patient to patient. Thus, pain is a highly subjective sensation and due to this fact, it becomes very difficult to quantify it in scientific researches⁽¹⁰⁾.

In many studies, Visual Analogue Scale (VAS) was verified to be sensitive to treatment effects⁽¹⁴⁻¹⁶⁾. It was found to correlate favorably with other self-reporting indicators of pain severity⁽¹⁴⁾. Furthermore, the difference in pain severity measured by VAS at two different points of time reflects the actual difference in pain level, which seems to be the main benefit of this method compared to others⁽¹⁷⁾.

Normally, pain during orthodontic treatment is noticeable, mainly in the first three days, reaching its maximum level in 24 hours, and decreasing after the third day of activation⁽¹⁰⁾.

According to the current results, the null hypothesis of the study was rejected. The present study employed a wavelength of 653 nm, power output 20 mW, energy density 6.5 J/cm². However, the outcomes obtained from the present study, demonstrated that there is a non-significant decrease in pain between the nonlaser and laser groups. This is in accordance with several previous studies^(3, 18-23), for example,Abtahi et al, AlSayed et al, Furquim et al, Lim et al, Angelieri et al, Dalaie et al, Heraviet al^(3, 18-23).

On the other hand, LLLT approved a positive effect toward reduction of pain perception during OTM ^(7, 13, 24-35). When Artés-Ribas et al, Nóbrega et al, Almallah et al, used a wavelength of 830 nm in conjunction with elastomeric separators placement^(24, 25, 30), it was effective to control pain experienced, which comes against the current study

findings. This could be due to the use of a higher wavelength. In addition, the higher dosage $12J^{(25)}$, $5J^{(30)}$, and higher energy density $16J/cm^{2(24)}$, could play a role in such dissimilar results.

After first and final archwires placement Tortamano et al, and Dominguez et al, found that LLLT with the same wavelength of (830nm) had a positive effect in pain reduction when used with energy density 5J/cm² per tooth for 16 s/point⁽³³⁾, and energy 4.4J per tooth,⁽³⁴⁾. These findings come against the current study, which could be due to the final higher dose delivered to each tooth.

However, Eslamian et al⁽²⁷⁾, used a wavelength of 810 nm and energy density 2J/cm² per point after elastomeric separators placemen, Farias et al⁽²⁸⁾ used the same parameters but in the early stages of orthodontic treatment. while Bayani et al⁽³¹⁾ used the same wavelength in the early stages of orthodontic treatment in conjunction with higher energy density 3.6J/cm². All of them found a significant decrease in pain perception after the LLLT application.

The controversy continued, when Qamruddin et al⁽¹³⁾, proved the positive effect of LLLT in pain reduction when used with higher parameters and doses than the current study. It was evident when applied a wavelength of 940 nm with a total energy of 12 J per tooth after elastomeric separation placement,⁽⁷⁾. in addition, when applied the same wavelength in combination with energy density of 75J/cm² per tooth in a single dose after every archwire placement during leveling and alignment stage⁽¹³⁾.

Marini et al⁽²⁹⁾, Deshpande et al⁽³²⁾, Bicakci et al⁽²⁶⁾, Doshi-Mehta, and Bhad-Patil⁽³⁵⁾ reduced the pain sensation with the help of LLLT. They used higher wavelengths of 910nm with a total energy of 54.4J per tooth after elastomeric separation^{(29),} 904nm with output 10 mW for 120 s/side during leveling and alignment phase^{(32),} 820nm with total energy 1J per tooth after elastomeric separators placement⁽²⁶⁾ and 800nm with 8 J per tooth after activation during canine retraction,⁽³⁵⁾ respectively.

CONCLUSION

Based on the current results, it could be concluded that the LLLT, with current parameters and protocol, has a negligible effect on pain experienced during initial orthodontic leveling and alignment stage.

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