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Clinical Assessment of the Efficiency of Low Level Laser Therapy in the Treatment of Oral Lichen Planus

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Abstract

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Key words: Diode laser; visual analogue scale; Oral lichen planus (OLP).

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Competing Interests: The authors have declared that no competing interests exist.

BACKGROUND: Oral lichen planus (OLP) is a chronic inflammatory disease of the oral mucosa of uncertain etiology.

AIM: To evaluate the effect of using low level laser therapy (LLLT (970 nm Siro laser Advance) for the treatment of symptomatic (OLP).

SUBJECTS AND METHODS: The present study was conducted on ten patients suffering from persistent oral lichen planus (OLP).Patients were treated with diode laser (970nm) for the symptomatic relief of pain and burning sensation. The patients were assessed before, during and after the completion of the laser treatment which was done twice weekly for two successive months with maximum of ten sessions. The assessment was performed using visual analogue scale (VAS) and clinical investigation for each patient.

RESULTS: Detailed significant reduction in lesion size and showed complete remission of burning sensation and pain. No reported complications or therapy side effects were observed in any of the treated patients.

CONCLUSION: Diode laser therapy seems to be an effective adjunctive treatment modality for relieving pain and clinical symptoms of OLP.

Introduction

Oral lichen planus (OLP) is a relatively common chronic inflammatory disease occurs in skin and oral mucosa of unknown etiology, it rarely undergoes spontaneous remission and potentially premalignant [1]. OLP mostly affects women between fifth and sixth decade. Approximately, (18-15%) of OLP patients are smokers and (24-29%) uses alcoholic beverages [2]. Clinical subtypes have been described. The white form include reticular, popular, and plaque-like forms while the red forms include atrophic (erythematous), erosive (ulcerative) and bullous [3]. The atrophic and erosive lesions are often associated with discomfort and other symptoms [4]. The pathogenesis involves a cell-mediated immune response to antigenic changes in oral mucosa; with prevalently patterns of inflammation described the plague, and erosive type. The plague type occurs most commonly and is estimated to affect up to 2% of population. It is present as purple raised areas with a white network known as Wickham's striae [6]. Several empirical treatments have been used including corticosteroids, curcuminoid, oxypentifylline, as well as surgery, photochemotherapy and lasers [7]. Low level laser therapy (LLLT) recently was used for treating erosive OLP with minimal side effects [8]. LLLT is an innovative approach in medicine, which has potential biostimulating effects when applied to oral tissues, improving wound healing [9] enhances epithelization after periodontal surgery [10]. Diode lasers provide great benefits over many other lasers because of its small size comparable to other types of laser. Diode laser also provides a wide range of spectrum that may be used in medical fields ranging

the predominantly T-lymphocytes infiltration [5]. Two

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from physiotherapy, photodynamic therapy, and surgical uses; it is transmitted through fibrotic [11]. The physical benefit used effectively in coagulation of superficial and interstitial lesions [12]. Several low level lasers used to treat oral lichen planus, including ultraviolet waves below (350 nm length), helium-neon, more recently, the diode laser. These lasers used with different wavelengths, intensities powers, durations, number of sessions and therapeutic approaches [13].

The aim of this study was to evaluate the effect of using low level laser therapy (LLLT (970 nm Siro laser Advance) for the treatment of symptomatic (OLP).

Subjects and Methods

The ethics committee approved the study protocol. The patients were collected from Department of Oral Medicine at the Faculty of Oral and Dental Medicine Cairo University. Ten patients with symptomatic (OLP) ranged in age from (45 to 60 years) unresponsive to topical steroids, were recruited from Dermatologic Department in Al-Qasr El Einy Hospital then referred to the Dental Clinic in National Research Center for laser treatment. Diagnosis of (OLP) is evaluated by clinical and histological features. Criteria of (OLP) diagnosis were established by World Health Organization (WHO) [14]. However, histopathological evaluation is recommended to verify clinical diagnosis and to exclude dysplasia and malignancy [15]. The selected patients had no known allergies or systemic diseases. Optimization of oral hygiene is fundamental to treat (OLP) with complete removal of supra and sub gingival plaque, whereas dental plaque and calculus implement intraoral inflammation and intensify both extensions and symptoms of (OLP) lesions. Patients were instructed to clean their teeth using a soft bristle toothbrush and toothpaste without irritants as mint or cinnamon, avoid accidental trauma on soft tissues; daily use of alcoholfree chlorhexidine mouth wash to reduce bacterial plaque, and prevent opportunistic fungal infection, especially in patients with dentures [16] throughout time of the study. Accidental toothbrush, trauma, pointed cusps, cracked teeth or worn dental restorations may worsen symptoms or may even trigger new lesions. Acidic, spicy, hard/crunchy and hot foods and beverages are not tolerated during active phases [17]. Patients with systemic disease or pregnancy were excluded. Patients were evaluated for demographics, medical history, presence of pain and discomfort, duration of disease, type, size, sites of oral involvement, these data were assessed and recorded. Patients were treated with diode laser (LLLT) 970 nm wavelength Sirolaser advance Germany). In a continuous non-contact mode with 320 µm diameter

fiber optic as delivery system that directed at the affected areas of oral mucous membrane with defocused mode and overlapping exposure until blanching of the treated area was observed. The patients and the operating staff were advised to wear special googles for eye protection. Diode laser was calibrated and applied with an output power of (3W), frequency at (30 Hz), energy at (180 joule) and time interval of eight minutes divided into four subsequent sessions each of two minutes and one minute rest in between to allow for tissue relaxation. A visual analogue scale (VAS) was used to rank the severity of the patient's symptoms ranging from zero (no pain) to ten(extreme pain) [18] and recorded before and after treatment. Clinical evaluation for the lesions was assessed and changes in the lesion's size were monitored at the respective laser sessions. Patients were trained and motivated to maintain proper plaque control, which included teeth brushing three times a day with a soft brush and tooth paste, and rinsing once daily with alcohol-free chlorhexidine mouth wash. After each laser session, a cold diet was recommended, whereas hot and spicy food was avoided.

Results

Changes within each week

Pain: At week one, there was no statistically significant change in pain scores through all week days.

At week two, there was no statistically significant change in pain scores from Day one to Day two. From Day two to Day three, there was a statistically significant decrease in pain scores. From Day three to Day four and through the rest of week days, there was no statistically significant change in pain scores.

At week three, there was a statistically significant decrease in pain scores from Day one to Day two. From Day two to Day three and through the rest of week days, there was no statistically significant change in pain scores.

At week four, there was no statistically significant change in pain scores from Day one to Day two and from Day two to Day three. From Day three to Day four and through the rest of week days, there was no statistically significant change in pain scores.

At weeks five, six as well as seven, there was a statistically significant decrease in pain scores from Day one to Day two. From Day two to Day three and through the rest of week days, there was no statistically significant change in pain scores. While at week eight, there was no statistically significant change in pain scores through all week days.

Table 1: Mean, standard deviation (SD) values of pain scores at different days within each week

Week	Day 1	Day 2	Day 3	Day 4	Day 5	Day 6	Day 7	P-value
	Mean	SD Mean	SD Mean	SD Mean	SD Mean	SD Mean	SD Mean	SD P-value
Week 1		2.0 7.1				2.1 7.0		
Week 2	6.8 ^a	1.9 6.7 ^a	1.9 6.4 ^b	2.1 0.019*				
Week 3	6.3 ª	2.2 5.9 ^b	2.1 5.9 ^b	2.1 5.9 ^b	2.1 5.8 ^b	2.0 5.8 ^b	2.0 5.8 ^b	2.0 0.001*
		2.1 5.0 ^b						
Week 5		2.1 4.0 ^b						
Week 6	3.7 ª	2.2 3.2 ^b	2.0 0.001*					
Week 7	2.9 ª	2.1 2.4 ^b	2.1 2.3 ^b	2.0 < 0.001*				
Week 8	2.1	2.1 2.1	2.1 2.1	2.0 2.0	2.2 2.0	2.1 2.0	2.1 1.9	2.0 0.911
*: Sianif	ficant i	atP≤0	.05. Differ	rent supe	rscripts in	the sam	ne row a	re statistically

significantly different.

Changes through different weeks

There was a statistically significant decrease in pain scores through all weeks.

Table 2: Mean, standard deviation (SD) values of pain scores at different days within each week

	Mean	SD	P-value	
Week 1	7.0 ^a	2.0		
Week 2	6.5 ^b	2.1		
Week 3	5.9 ^c	2.1		
Week 4	4.8 ^d	2.0	0.004*	
Week 5	4.0 ^e	2.1	<0.001*	
Week 6	3.3 [†]	2.1		
Week 7	2.4 ^g	2.0		
Week 8	2.0 ^h	2.1		
*: Cignificant of	R < 0.05 Different our	orcorinto in the	anno column ara a	totiotioolly

*: Significant at $P \le 0.05$, Different superscripts in the same column are statistically significantly different.



Figure 1: Line chart representing mean pain scores at different weeks

Lesion area

There was a statistically significant decrease in mean lesion area after treatment. The lesion area decreased by $90.0 \pm 10.6\%$ after treatment.

Table 3: Mean, standard deviation (SD) values and results of comparison between lesion area before and after treatment

Before treatment		After trea	After treatment		ase	P-value
Mean	SD	Mean	SD	Mean	SD	
318.2	326.2	46.0	102.8	90.0	10.6	<0.001*
*: Significant at P ≤ 0.05						

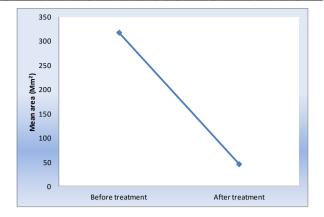


Figure 2: Line chart representing mean lesion area before and after treatment

Initial improvement of OLP started after two weeks of treatment including the decrease in pain. Fig 1 show statistical significant decrease in pain scores starting from day two in the second week reaching its minimum score by week eight, but the patient continued to complain pain in the region of the lesion stimulated by hot and spicy food. Clinical remission was obtained in a period of two months and complete disappearance of the lesion in a period of two months is shown in Fig. 2. In this study the pain score show a high statistical improvement. Most patients reported an immediate pain relief after the second session, and all of them reported a complete resolution of symptoms at the end of the laser session, even if some lesions show partial clinical response. The results are concomitant with (Hearty, 2002) study which reported that the quality of life of patients with (OLP) improved significantly with the absence of complete resolution of all oral signs with no reported complications [19]. Cafaro et al., [20] treated 13 patients with (OLP) using the pulsed diode laser, he concluded that there was a significant decrease in lesions and decreased pain without any side effects.

Discussion

Oral lichen planus (OLP) is a chronic inflammatory autoimmune disease characterized by remission and exacerbation. It is a cell-mediated immune condition. in which T-lymphocytes accumulate beneath the epithelium of the oral mucosa and increase rate of differentiation of stratified squamous epithelium, resulting in hyperkeratosis and erythema with or without ulceration [21]. T-cells (mostly CD8+ and CD4+ cells) migrate into the epithelium due to random encounter of antigen during routine surveillance [22]. The migrated (CD8+) cells are activated directly by antigen binding to major histocompatibility complex (MHC-1) on keratinocyte or activated (CD4+) lymphocytes [23]. The activated T-

cells and amplified (Th1) cytokine production (IL-1, IL-8, IL-10, IL-12, tumor necrosis factor-alpha (TNF- α) enhancing the expression of intercellular adhesion molecule-1 (ICAM-1) on Langerhans cells and macrophages and major histocompatibility type 1 (MHC-1) antigen on keratinocytes [24]. Patients affected with erosive-atrophic lichen planus may exhibit extensive and painful lesions which are often conventional unresponsive to treatments. Corticosteroids are most widely accepted treatment for (OLP), which relieves symptoms rather than curing it [27]. Low level laser therapy (LLLT) is recent evolution on medical/dental treatments, specifically mucocutaneous lesions as (OLP) [25]. The primary benefit of (LLLT) being non-surgical, it promotes tissue healing and reduce edema, inflammation and pain. Its principle of using (LLLT) is to supply direct biostimulation light energy to body's cells. LLLT accelerate healing, anti-inflammatory effects, increase cellular metabolism, modulation of immune system, vasodilatation and analgesic effects, and amelioration of morphine withdrawal [26]. Other modalities like calcineurin inhibitors, retinoids, dapsone, hydroxychloroiquine, and enoxaparin have contributed significantly towards the disease. The pathogenesis of the disease suggests that blocking (IL-12), (TNF- α), RANTES or (MMP-9) activity or upregulation (TGF-β1) activity on (OLP) may be therapeutic value in future [28]. The principle of using LLLT is to supply direct biostimulation light energy to body's cells. Diode laser (980 nm) possesses a deep power of penetration reaching about 1.5 mm [29]. Application of diode (980 nm) will rise in temperature of affected tissues to above 50 degrees and less than 100 degrees causing protein denaturation causing blanching of treated mucosa; this denaturation means the destruction of the diseased epithelium with its surface antigen [30]. Van der Hem et al., [31] reported that re-epithelizaton occurs within three weeks after removal of the epithelium by the laser and feeling discomfort when in contact with food or liquid disappears. All laser types destroy the superficial epithelium containing target keratinocytes by protein denaturation [32]. Our results showed that the use of LLLT in the management of OLP has a significant effect on reducing pain and clinical parameters of inflammation. The initial improvement of atrophic-erosive lesions started after two weeks of treatment. Clinical remission was obtained in a period of approximately one month. Lesions that do not show any clinical improvement after four laser sessions were excluded from the study, this means that type of patient does not respond clinically, and the treatment should be stopped. The pain score had a high statistical improvement and patients successfully reported a great enhancement in quality of life with complete resolution of symptoms at the end of laser sessions. LLLT act analoesically since they improve endorphin release and therefore inhibit nociceptive signals and control pain mediators [33]. The size, site, depth, seemed to influence healing. The small number of

patients diminished the possibility to draw any conclusions with statistical significance about parameters that might had more influence on the results, so further investigations with other treatment modalities and a prolonged follow-up to confirm the efficacy of laser therapy in the treatment of symptomatic OLP.

References

1. Carbone M, Arduino PG, Carrozo M, et al. Course of oral lichen planus a retrospective study. Italian patients. Oral Dis.2009;15:235-243.

http://dx.doi.org/10.1111/j.1601-0825.2009.01516.x PMid:19222766

2. Oliveria MG, Almeida JD, Balducci I. Study of 110 Brazilian patients. BMC Res Notes. 2010;3:157. http://dx.doi.org/10.1186/1756-0500-3-157 PMid:20525297 PMCid:PMC2898663

3. Ismail SB, Kumar SK, Zain RB. Oral lichen planus and Lichenoid reaction: Etiopathogenesis, diagnosis, management and malignant transformation. J Oral Sci. 2007;106:49-89. http://dx.doi.org/10.2334/josnusd.49.89

4. Crincoli V, Di Biscelgie MB, Scivetti M, Festa F. Oral lichen planus: update on etiopathogenesis, diagnosis and treatment. Immunopharmacol Immunotoxicol. 2011;33:11-20. <u>http://dx.doi.org/10.3109/08923973.2010.498014</u> PMid:20604639

5. Brightman V. Red and white lesions of oral mucosa. In Burkit's Oral Medicine, diagnosis and treatment.10th Edition, edit by Greenberg M, and Glick M, 2003.

6. Carrozo M. How common is oral lichen planus? Evidence-Based Dentistry. 2008;9(4):112-3.

http://dx.doi.org/10.1038/sj.ebd.6400614 PMid:19151683

7. Conrotto D, Carbone M, Carrozo M, Arduino P, Broccoletti R, Pentenero M, et al. Ciclosporin vs Clobetasol in the topical management of atrophic and erosive oral lichen planus: a doubleblind, randomized controlled trial. Br J Dermatol. 2006;154:139-145.

http://dx.doi.org/10.1111/j.1365-2133.2005.06920.x PMid:16403107

8. Passeron T, Zakaria W, Ostovari N, Mantoux F, Lacour JP, Ortonne JP. Treatment of erosive oral lichen planus by 308 nm excimer laser. Surg Med. 2004;34:205. http://dx.doi.org/10.1002/lsm.20016 PMid:15022246

9. Gal P, Vidinsky B, Toporcer T, et al. Histological assessment of the effect of laser irradiation on skin wound healing in rats. Photomed.Laser Surg. 2006;24:480-484. http://dx.doi.org/10.1089/pho.2006.24.480 PMid:16942428

10. Ozcelik O, Cenk Haytac M, Kunin A, Seydaoglu G. Improved wound healing by low-level laser irradiation after gingivectomy operations: a controlled clinical pilot study. J Clin Periodontol. 2008;35(3):250-4.

http://dx.doi.org/10.1111/j.1600-051X.2007.01194.x PMid:18269665

11. Moritz A, Gutknecht N, Doertbudak O, Goharkhay K, Schoop U, Schauer P, Sperr W. Bacterial reduction in periodontal pockets through irradiation with diode laser, a pilot study. J Clan Laser Med Surg. 1997;15:33-37. PMid:9467340

12. Romanos G, Nentwig GH. Diode laser (980 nm) in oral and maxillofacial surgical procedures: clinical observations based on

clinical applications. J Clin Laser Med Surg. 1999;17(5):193-7. PMid:11199822

13. Jajarm HH, Falaki F, Mahdavi O. A comparative pilot study of low intensity laser versus topical corticosteroids in the treatment of erosive-atrophic oral lichen planus. Photomed Laser Surg. 2011;29:421-425.

http://dx.doi.org/10.1089/pho.2010.2876 PMid:21214369

14. Rad M, Hashemipoor MA, Mojtahedi A, Zarei MR, Chamani G, Kakoei S, Izadi N. Correlation between clinical and histopathologic diagnoses of oral lichen planus based on modified WHO diagnostic criteria. Oral Surg Oral Med Oral Pathol Oral Radiol Endod. 2009;107(6):796-800.

http://dx.doi.org/10.1016/j.tripleo.2009.02.020 PMid:19464654

15. Scully C, Carrozo M. Oral mucosal disease: Lichen planus. Br J Oral Maxillofac Surg. 2008; 46:15-21. http://dx.doi.org/10.1016/j.bjoms.2007.07.199 PMid:17822813

16. Salerno C, Pascale M, Contaldo M, Esposito V, Busciolano M, Milillo L, Guida A, Petruzzi M, Serpico R. Candida-associated denture stomatitis. Med Oral Patol Oral Cir Bucal. 2011;16(2):e139-43. http://dx.doi.org/10.4317/medoral.16.e139

http://dx.doi.org/10.4317/medoral.16.e139 PMid:20711156

17. Schlosser BJ. Lichen planus and lichenoid reactions of the oral mucosa. Dermatol Ther. 2010;23:2.

http://dx.doi.org/10.1111/j.1529-8019.2010.01322.x PMid:20597944

18. Scott J, Huskisson EC. Graphic representation of pain. Pain. 1976;2(2):175-84.

http://dx.doi.org/10.1016/0304-3959(76)90113-5

19. Hegarty AM, Hodgson TA, Lewsey JD, Porter SR. Fluticasone propionate spray and betamethasone sodium phosphate mouthrinse:a randomized crossover study for the treatment of symptomatic oral lichen planus. J Am Dermatol. 2002;47:271-279. http://dx.doi.org/10.1067/mjd.2002.120922

20. Cafaro A,Albanese G,Arduino PG,Broccoletti R(2010).Effect of low-level laser irradiation on unresponsive oral lichen planus: early preliminary results in 13 patients.Photomed Laser Surg;28 suppl 2:s99-103.

http://dx.doi.org/10.1089/pho.2009.2655 PMid:20738169

21. Dissemond J. Oral lichen planus: an overview Dermatology Treat. 2004;15:136-40.

22. Zhou XJ, Sugerman PB, Savage NW, Walsh LJ, Seymour GJ. Intra-epithelial CD8+ T cells and basement membrane disruption in oral lichen planus. J Oral Pathol Med. 2002;31(1):23-7. http://dx.doi.org/10.1046/j.0904-2512.2001.10063.x PMid:11896819

23. Zhou XJ, Sugerman PB, Savage NW, Walsh LJ, Seymour GJ. Intra-epithelial CD8+ T cells and basement membrane disruption in oral lichen planus. J Oral Pathol Med. 2002;31(1):23-7. <u>http://dx.doi.org/10.1046/j.0904-2512.2001.10063.x</u> PMid:11896819

24. Scardina GA, Ruggieri A, Messina P, Maresi E. Angiogenesis of oral lichen planus: a possible pathogenetic mechanism. Med Oral Patol Oral Cir Bucal. 2009;14(11):e558-62. http://dx.doi.org/10.4317/medoral.14.e558 PMid:19680199

25. Jajarm HH, Falaki F, Mahadavi O. A comparative pilot study of low intensity laser versus topical corticosteroids in the treatment of erosive-atrophic OLP. Photomed Laser Surg. 2011;29:421-425. http://dx.doi.org/10.1089/pho.2010.2876 PMid:21214369

26. Mirzaii-Dizgah I, Ojaghi R, Karimian SM, Sohanaki H. Attenuation of morphine withdrawl signs of low level laser therapy in rats. Behav Brain Res. 2009;196(2):268-70. <u>http://dx.doi.org/10.1016/j.bbr.2008.09.015</u> PMid:18926854 27. Sharma S, Saimbi CS, Koirala B. Erosive oral lichen planus and its management: a case series. J Nepal Med Assoc. 2008;47(170):86-90. PMid:18709038

28. Roopashree MR, Gondhalekar RV, Shukla A. Pathogenesis of oral lichen planus-a review. J Oral Pathol Med. 2010; 39:729-34. http://dx.doi.org/10.1111/j.1600-0714.2010.00946.x PMid:20923445

29. Captone GA, Halusic E. Photobiology of laser in oral maxillofacial surgery. In laser application in oral & maxillofacial surgery. Catone GA and Alling C. Saunder company, 1997:29-38.

30. L'Esperance FA. Ophthalmic laser. Third edition, chapter 1, 1989:1-10.

31. van der Hem PS, Egges M, van der Wal JE, Roodenburg JL. CO2 laser evaporation of oral lichen planus. Int J Oral Maxillofac Surg. 2008;37(7):630-3. http://dx.doi.org/10.1016/j.ijom.2008.04.011

PMid:18538542

32. Trehan M, Taylor CR. Low-dose excimer 308-nm laser for the treatment of oral lichen planus. Arch Dermatol. 2004; 140:415-20. http://dx.doi.org/10.1001/archderm.140.4.415 PMid:15096369

33. Tezel A, Kara C, Balkaya V, Orbak R. An evaluation of different treatments for recurrent aphthous stomatitis and patient perceptions: Nd:YAG laser versus medication. Photomed Laser Surg. 2009;27(1):101-6. http://dx.doi.org/10.1089/pho.2008.2274

PMid:18687056