The Use of Soft Tissue Diode Laser in the Treatment of Oral Hyper Pigmentation

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Abstract

Objective: To evaluate the clinical effectiveness and patients’ acceptance of soft tissue diode laser, for the elimination of hyper pigmented areas of the gingivae and buccal mucosa.

Subjects and methods: Twenty patients complaining of dark localized areas in their gingivae of upper and lower jaws, or in cheek mucosa were included in this study. All are females ages ranged from 18-37 with average of 27.5 years. The patients were all not pregnant, and free from any systemic diseases, and had no other areas of hyperpigmentation.

Results: Most of the patients (18 Pts.) required two treatment sessions to remove the melanotic lesions satisfactorily. The patients were seen after one, two, and three weeks respectively. Eight patients (40%) had a good response after one week, Ten patients (50%) had a very good response after two weeks, while only two patients (10%) had a bad result and required a third treatment session after two weeks. At the third week postoperatively in the successful cases, restoration of the normal gingival and mucosal texture and color was achieved with optimal esthetic results. Most of the patients had no postoperative discomfort including: pain, edema and disfigurement.

Conclusion: The soft tissue diode laser can be considered as a reliable mean for elimination of melanotic lesions of the gingivae or buccal mucosa, and is well tolerated by the patients.

Key words: hyperpigmentation- soft tissue diode laser- patient response

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Introduction:

Oral pigmentation can appear at any age in the gingivae, mucosa, hard palate and tongue. These can be of genetic origin as racial or physiological pigmentation, Pigmented nevi, and Albright syndrome. Acquired oral pigmentation can occur as a result of endogenous or exogenous factors. The endogenous pigmentation often occurs during the course of certain diseases like Addison’s disease, Acanthosis Nigricans, and Cooley’s anemia, while exogenous pigmentation may be due to amalgam tattoo, lead poisoning, Argyria, antimalarial drugs or smoker’s melanosis. (1) Hyper pigmentation of gingiva often occurs in the gingiva due to abnormal deposition of melanin in the basal and supra basal layer of the epithelium, creating a unaesthetic appearance. The effective elimination of such melanotic areas was attempted by variable means such as; scalpel surgery; gingivectomy with free-gingival autografting or allografting, (2,3) electrocautery, cryosurgical depigmentation through the use of gas expansion system, (4) chemical agents such as phenol 90% and alcohol 95%, (5) abrasion with diamond burs (3) as well as laser surgery such as; Neodymium -Yttrium garnet (Nd:YAG) Laser, Co₂ Laser and semi-conductor diode Laser. (6) Researchers in selective photothermolysis and treatment of cutaneous pigmented lesions have been performed with various lasers. (7,8) Selective photothermolysis to treat melanotic lesions was first proposed by Anderson and Parrish in 1983. (9) This phenomenon is based on the fact that some structures absorb light of certain wavelength better than others. The absorption of light leads to heat production and consequently to damage of structures when a high enough temperature is reached. As absorption by each structure is wavelength dependent, it is possible to selectively damage certain regions in a tissue by choosing a wavelength with high absorption by the target and low absorption by others. (2,9) Although the literature reports great variability in the values, (3,10) the absorption coefficient of melanosomes for red light may be estimated to be several hundreds of reciprocals centimeters. In addition to the wavelength the proper pulse length is another parameter that is essential for the achievement of selective photoagulation decreases the selectivity of tissues to photothermolysis. Research in selective photothermolysis and treatment of pigmented lesions has been accomplished with various lasers, including 10.600 nm Co₂, (2) 1060nm, 532 nm and 355 nm Neodymium–Yttrium garnet 351 nm excimer, (6) 694 nm Quartz-switched ruby (7) and 488/514 nm argon laser. (7,9,10) Most of these are pulsed lasers that are able to produce pulses smaller than, or in the order of magnitude of, the thermal relaxation time of a melanosome which is around 1 microsecond. The soft tissue Diode laser SoftLase Pro is a compact and reliable surgical instrument for use in dental applications. Flexible fiber optic cables are the most widely accepted delivery system for diode laser power; the 400μm diameter single-core quartz fiber is mounted into standard connectors. It produces an invisible infra-red and a visible aiming beam. The wavelength is 808±5nm, such wavelength characteristics specially target the soft tissues. It has an affinity for hemoglobin and melanin; therefore it is more efficient and better equipped to address deeper soft tissue problems. (5,11)

Aim of the work: The purpose of this study was to assess the clinical effectiveness and patients’ responsiveness for the Soft Tissue Diode laser in the elimination of gingival and mucosal hyperpigmentation.

Subjects and Methods:

Twenty patients complaining of unsightly hyper pigmented areas in their gingivae, labial and cheek mucosa were included in this study, especially those patients with gummy smile (very high gum line). The patients were all females, their ages ranged from 18-37 with average of 27.5 years, from those attending the Outpatient Dental Clinic of Taif University Girl’s branch. Clinical assessment was made prior to treatment at the first visit. All patients were non pregnant, they were apparently healthy without any systemic diseases, or manifestations of other areas of hyper pigmations, as nail pigmentation–etc. All patients were non-smokers and were not exposed to metal vapors as occupational hazards i.e. the melanotic spots were of physiological or racial nature. The procedure was explained verbally and animatedly to the patients and informed signed consents were
obtained according to the World Declaration Statement of Helsinki.\(^{12}\)

**Methods (Treatment technique):** Patient and operating staff wore special diode-laser protective eye glasses. Highly reflective instruments or instruments with mirrored surfaces were avoided as there could be reflection of the laser beam. A soft tissue diode surgical laser unit (Discuss SoftLase Pro, LLC of ZAP Lasers, Culver City, CA 90232 USA), delivered by a 400-μm diameter fiber wavelength 808nm ± 5, power 1-2 W was used for depigmentation of the maxillary and mandibular anterior gingiva and buccal mucosa. The diode laser was emitted in continuous mode, and was operated in a contact method using a flexible fiber optic hand piece with initiated tip. The Laser settings used were: fluent between 5 and 15 J/cm², pulse length between 20 and 50 ms, spot diameter of 2 mm and thermal relaxation times of 0.15 to 0.20 milliseconds. No topical or local anesthesia was given to patients. Laser ablation started from the mucogingival junction toward the free gingival margin, including papillae. Regarding the labial or cheek mucosa the motion of ablation was circular with overlapping circles. During the procedure, laser ablated the gingival and or mucosal epithelial surfaces little by little to reach the pigments without causing any bleeding which was beneficial for clear visualization. To enhance visualization, normal saline-soaked cotton or gauze was used to remove epithelial remnant. The power output at the end of the fiber was adjusted with power increase or power decrease touch pad until it measured from 1.2W up till 2W. The used treatment was aiming beams to irradiate the pigmented lesions at 1.5 W. Pulse rate selected was with contact fiber mode (initiated tip). Anesthesia was not used in all cases, but if needed according to patient's desire; local anesthesia was achieved using 1.8 ml mepivacaine 2% without vasoconstrictor, although more than one carpile was used when large areas were treated. After about 3-5 minutes, the treatment began. Irradiation was done by a contact mode for the fiber; a slight degree of carbonization at the gingival or mucosal surface was seen. During treatment the temperature of the tissue often quickly reaches high values. It was not necessary to treat the entire surface of the lesion. In spite of the beam being focused to a spot size of only 2mm in diameter, the heat effect exceeded this diameter as a result of conduction. Irradiation of spots approximately 1mm apart appeared to sufficiently affect the area in-between. A thin layer of carbonized material appeared at the irradiated spot. After having irradiated the pigmented lesions, the wounds appeared fresh with no bleeding and there were no need to apply periodontal dressing or pack. The patients were instructed to avoid eating hot or spicy food for the first 24 hours and to gargle with Aloe Dent mouth wash 3 times/a day preferably after meals for one week. No pain medications were generally prescribed, but if needed a mild analgesic anti-inflammatory (sodium diclofenac 50mg =Rofenac) twice a day was given.

**Post-operative assessment:** The patients were seen after one, two, and three weeks post-operatively, and were examined for post-operative discomfort; including pain, edema, and disfigurement in the form of hypopigmentation (i.e. treatment response). Clinical parameters were measured using Visual Analogue Scale (VAS), through answering a questionnaire specifically designed for this research.\(^{13}\)

**Results:**

No patients had marked complications, and infection did not occur during the healing of the gingiva. In two patients (10%), some areas of pigmentation remained and required a third session after two weeks. Regarding pain: it was assessed using Visual Analogue Scale.\(^{13,14}\) Operationally a VAS is usually a horizontal line, 100 mm in length, anchored by word descriptors at each end (no pain at one end till very severe pain on the other end). Patients were asked to mark on the line the point that they feel represents their perception of their current state. The VAS score is determined by measuring in millimeters from the left hand end of the line to the point that the patient marks.

Only three patients (15%) marked on a point equivalent to mild pain and thus they required mild analgesics. Regarding edema: It was measured by a graduated tape \(^{14}\) although it was not seen in most of the patients at the first week, only two patients(10%) presented with slight edema in the attached
gingiva at the mucogingival junction, which disappeared completely at the 2nd week. While in patients with cheek mucosal depigmentation no post ablation edema was seen.

Regarding the post-operative appearance and the patient acceptance for the treatment (treatment response): It was registered on a scale measuring patients' satisfaction with the present situation and the psychologic impact of their oral health status with their responses marked on a Visual Analog Scale (VAS) with four values subjectively (very good, good, fair, and bad). A very good score was registered when both the dentist and the patient were satisfied with the final result. A good response was considered to be achieved when the solution was satisfied by the result, but the patient considered that it would have been better. A fair result was scored when both the doctor and the patient estimated that the treatment could have given a better result. A bad result (no response) was registered when the treatment did not produce noticeable positive changes from the initial status.

Results were graphed as seen in Figure (1).

**At the First post-operative week:**
Eight patients (40%) were classified as having a good result. Ten (50%) as having a fair result, while only two patients (10%) as having a bad result i.e. no response.

**At the Second week post-operatively:**
Twelve patients representing (60%) were seen with a very good result, three patients (15%) with good result and another three patients (15%) had fair result, and two patients (10%) came with a bad result.

**At the third post-operative week:**
Thirteen patients representing (65%) were seen with a very good results, while six patients (30%) came with good results, and only one patient (5%) presented with a fair result. More than half of the patients (12pts.) included in the study were available within nine months follow-up, during this period no recurrence of the pigmentation appeared, while the remaining patients did not show up.
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Fig (5): showing the application of soft tissue diode Laser beam.

Fig (6): Immediate post-operative photo showing slight reduction in gingival hyperpigmentation in upper rather than the lower gingiva.

Fig (7): Post-operative photo after application of Laser for upper gingiva only. Notice the difference between upper and lower untreated (one week postoperatively).

Fig (8): Preoperative photo showing hyperpigmentation in the cheek mucosa.

Fig (9): Immediate postoperative photo showing the effect of laser beams on the cheek mucosa.

Fig (10): Two weeks postoperatively showing the improvement of the dark appearance of the cheek mucosa.
Discussion

Although the melanin pigmentation of the gingiva is a completely benign condition and does not present a medical problem, yet complaints of black gums are common. Demand for treatment is usually made for esthetic reasons however; there is not much information in the literature about the depigmentation of the gingiva.\(^7\)

Several treatment modalities were applied for the treatment of such melanotic spots Perlmutter et al,\(^7, 15\) and Dummett and Bolder\(^16\) have reported surgical removal of portions of pigmented gingiva. After surgery, it was necessary to cover the exposed lamina propria with periodontal packs for 7 to 10 days; the wound took 6 weeks to heal and left a delicate scar.

Cryosurgery is a method of tissue destruction by rapid freezing. There have been several reports on the cryosurgical treatment of a variety of oral lesions\(^15, 16\) but only Tal et al,\(^10\) described the depigmentation of gingival by cryosurgery. Others reported the successful use of liquid nitrogen and cotton swabs to eliminate the pigmented areas.\(^3\)

Various lasers have been utilized for the treatment of cutaneous pigmented lesions.\(^17, 18, 19\) Trelles et al,\(^19, 20\) were the first to treat patients with pigmented gingiva by Argon laser. The Ruby laser has been successfully used in Q-switched mode to treat a variety of pigmented lesions.\(^3, 6, 11\) The short pulses of the light emitted in the nanosecond domain by the Q-switched Ruby laser produce thermal and or mechanical damage to melanosomes, which results in cellular destruction confirmed to the targeted melanocyte, a process, which has been termed selective photothermolysis. Regarding the post-operative clinical assessment of pain and edema, diode laser was well tolerated by the patients where analgesics were not prescribed only if necessary, in accordance with other investigations,\(^21, 22\) as well as edema was minimally found and in limited cases.\(^23, 24\)

In agreement with this study, several studies have shown melanosomes to be suitable targets for selective photothermolysis at various laser wavelengths when irradiated in sub microsecond pulses with energy densities around1J/cm\(^2\).\(^9, 25, 26\) The pulse length is the important difference between the parameters as used in this clinical study for elimination of melanotic lesions, and those described in most of the literature of the pigmented lesions of the skin.\(^6, 9, 11\) Excellent results were achieved with pulse lengths of 300ms being much longer than the estimated relaxation time for the melanosomes. It is obvious that a mechanism different from ordinary photothermolysis plays a role in the elimination of the pigmented oral lesions using the described parameters. During the relatively long pulses, a considerable part of heat is conducted away from the targets to surrounding structures and general heating is the result.\(^9, 10\) Although a relatively low power is used (1.4W) in the present study, this in combination with a spot size of 0.5mm and a pulse length 300ms leads to a very high energy densities. As a result of both the heat diffusion and the light diffusion through scattering, two main effects appear from irradiation with the previously mentioned parameters; a coagulated and necrotized layer, together with an underlying area that is only partially coagulated, the extent of irreversibly damaged melanocytes gradually decreases with depth in this underlying area. This may be a crucial point in the mechanism of action,\(^24, 25\) although the remaining melanocytes may be fewer than normal, yet they may produce enough melanin to re-pigment the newly formed mucosa during and after the healing period, thus equalizing the pigmentation with the surrounding tissue.\(^16\) In the present study, it appeared that the melanocytes where not sufficiently destructed in all cases, so that repeated laser applications were necessary especially in the patients with fair and bad results. It was reported that, enough melanocytes are destroyed to remove the over pigmentation but, at the same time, some underlying melanocytes are not sufficiently affected by the heat to be destroyed.\(^26, 27\)

Some other mechanisms of re-pigmentation might be found in those areas in which melanocytes have been removed or destroyed. As for example migration of melanocytes several weeks after treatment. Although the 300ms pulses allowed the heat produced to partially damage the non melanin containing structures, the wounds healed with a good cosmetic appearance, this finding was previously supported in the literature.\(^19, 20\)

Treatment with short pulsed Lasers may be theoretically safe but those lasers are not
commonly available in hospitals or clinics. In contrast, the soft tissue diode laser is a common instrument, available in most clinics and hospitals due to its cheapness and lightness so it is easily transported from place to place. Although the CO\textsubscript{2} laser is preferred over the Argon laser and the Nd: YAG lasers, because it is most selectively absorbed, the CO\textsubscript{2} laser totally evaporates the tissue layer by layer, as its light is very well absorbed by any tissue containing water,\textsuperscript{(1,5,28)} while the Argon and Diode Lasers light share the same mechanism of action of the Nd:YAG laser which penetrates into the gingival tissue and is only absorbed by the melanin, resulting in much more selective destruction and far less damage to normal gingival tissue than does the CO\textsubscript{2} laser.\textsuperscript{(29,30,31)} In accordance to what previously mentioned in this present study, Bradley \textsuperscript{(37)} in a review for the use of the Nd:YAG Laser in oral and maxillofacial surgery, reported that Nd: YAG Laser wavelength exhibits minimal tissue absorption and maximal penetration by comparison with the CO\textsubscript{2} laser’s maximal temporo-mandibular joint. Promising developments include CO\textsubscript{2}; Nd: YAG lasers combination and the introduction of diode technology.\textsuperscript{(18)}

We conclude that in comparison with the treatment modalities discussed previously to treat the pigmented areas of the gums, the diode laser is a useful and a safe mean and is the preferred laser when no other short pulse Lasers are available.

**Conclusion:**

In conclusion, it is reported that the soft tissue diode laser wavelength exhibits maximal tissue absorption and minimal penetration by comparison with the Nd: YAG Laser’s minimal tissue absorption and maximal penetration.\textsuperscript{(1)} These properties allow a variety of uses in maxillofacial surgery particularly coagulation of angiomatous lesions, hemostasis in bleeding diatheses, arthroscopic surgery of temporo-mandibular joint. Promising developments include CO\textsubscript{2}; Nd: YAG combination Lasers, Potassium titanyl-phosphate (KTP); Nd: YAG apparatus and the introduction of Diode technology.\textsuperscript{(5)} Therefore, in comparison with the treatment modalities mentioned previously to treat the pigmented areas of the gums, the Diode Laser is a useful and a safe mean, and is the preferred laser when no other short pulse Lasers are available.

**References:**

Adopted by the 18th WMA General Assembly, Helsinki, Finland, June 1964, and amended by the 59th WMA General Assembly, Seoul, October 2008


