Published online 2015 December 29.

Case Report

Successful Treatment of Resistant Discoid Lupus Erythematous (DLE) With Low-Level Laser Therapy (LLLT): A Case Report

Nooshafarin Kazemikhoo,¹ and Parvin Mansouri^{1,*}

¹Skin and Stem Cell Research Center, Tehran University of Medical Sciences, Tehran, Iran

^{*} *Corresponding author*: Parvin Mansouri, Skin and Stem Cell Research Center, Tehran University of Medical Sciences, Tehran, Iran. Tel: +98-2122201710, E-mail: mansouripr@yahoo.com

Received 2015 September 15; Revised 2015 November 15; Accepted 2015 December 15.

Abstract

Introduction: Lupus is a chronic autoimmune illness characterized by production of autoantibodies against nuclear antigens. Lupus may also only appear as a skin disease. Discoid lupus erythematous (DLE), the most common form of chronic cutaneous lupus erythematous is characterized by patches of persistent erythema, adherent scaling, follicular plugging and, in the later stages scaring and atrophy.

Case Presentation: The case was a 33-year-old female with two years history of DLE, who referred to the laser clinic of Milad hospital with acute flare of skin lesions since seven months ego. She had four lesions (three with local alopecia, scaling and itching and one with atrophy). Serological studies for antinuclear antibodies and anti Ro/SS-A antibody were negative. After five sessions of treatment with infrared (830 nm; 200 mw) and red light (630 nm; 20 mw) diode laser in addition to intravenous red light laser (655 nm; 2 mw) for 20 minutes and laser acupuncture with infrared diode laser (1 J/cm²) for six acupuncture points every other day, the beneficial effects started to appear, and after nine sessions, local itching of the lesions had stopped. Through a total of 27 sessions of treatment, the lesions improved completely with highly satisfactory results; and drug-therapy held to be continued. No disease flare up was seen after the one-year follow up.

Conclusions: To the best of our knowledge, this is the first case report on low-level laser therapy (LLLT) treatment of DLE. Low-level laser therapy may be effective in treating inflammatory lesions such as cutaneous lesions of discoid lupus erythematous. More studies with higher sample size are needed to clarify the role of low level lasers in treating such lesions.

Keywords: Low Level Laser Therapy, Discoid Lupus Erythematous, Inflammatory Cutaneous Lesions, Laser Acupuncture, Intravenous Laser Therapy

1. Introduction

Discoid lupus erythematous (DLE), the most common form of chronic cutaneous lupus erythematous is characterized by patches of persistent erythema, adherent scaling, follicular plugging and, in later stages scaring and atrophy (1, 2).

These lesions persist for years and tend to expand slowly. Biopsy of DLE lesions shows hyperkeratosis, follicular plugging, and atrophy of the epidermis; hydropic degeneration of basal keratinocytes; and a mononuclear cell infiltrate adjacent to epidermal, adnexal, and micro vascular basement membranes (3). Treatment is focused on control of local cutaneous disease and consists mainly of photo protection and topical treatment with glucocorticoids, immune modulators including Elidel and Protopic, intra-lesional glucocorticoids and oral medications including antimalarials, Thalidomide and Dapsone. If local therapy is ineffective, use of aminoquinoline antimalarials may be indicated (4, 5).

High power lasers including PDL have been used for treatment of chronic refractory lesions of discoid lupus

erythematous since 1986 (6). We herein report, the first case of DLE treated with specific low-level laser therapy.

2. Case Prtesentation

The case was a 33-year-old female with six years history of DLE, who was referred to the laser clinic of Milad hospital (2009) with acute flare of skin lesions since seven months ego. She had four lesions (three with local alopecia, scaling and itching and one with atrophy) on the scalp (Figure 1A).

The histological examination of lesions supported the clinical diagnosis of DLE. Serological investigations for antinuclear antibodies, anti Ro/SS-A antibody, erythrocyte sedimentation rate (ESR), rheumatoid factor (RF) were negative. However, anti ds-DNA antibody was positive. The patient showed no signs of underlying systemic illness manifestations. Her skin lesions were under control with standard therapies including local corticosteroids and antimalarial drugs (hydroxychloroquine 200 mg/day) for several months, however after withholding antimalarial drugs because of pregnancy and breastfeeding period, the

Copyright © 2015, Skin and Stem Cell Journal. This is an open-access article distributed under the terms of the Creative Commons Attribution-NonCommercial 4.0 International License (http://creativecommons.org/licenses/by-nc/4.0/) which permits copy and redistribute the material just in noncommercial usages, provided the original work is properly cited.



Figure 1. Discoid Lesions on the Patient's Head A, Before and B, After Application of Low-Level Laser Irradiation; Note the Size of the Lesions Before and After Low-Level Laser Therapy

lesions flared up during and subsequent to pregnancy. Intra-lesional injections of corticosteroids (diluted triamcinolone acetate, 4 mg/mL) and antimalarial therapy were started. However, there was no response from skin lesions on the scalp during about seven months of corticosteroid oral therapy.

We used a combination of low-level laser therapy using local irradiation of lesions with infrared and red light diode laser (830 nm, 200 mw; and 630 nm, 20 mw, respectively; each 6I/cm²; Mustang 2000, Moscow, Russia) in addition to intravenous red light laser (655 nm, 2 mw for 20 minutes). Intravenous laser therapy was done using a fiber optic laser conduit inserted via a needle placed on the forearm in median cubital vein. We also used laser acupuncture with infrared diode laser (1 J/cm² Mustang 2000, Moscow, Russia) for LI-11, LI-4, SP-6, PC-6, ST-36 and GB-34 points. All three methods of laser therapy (local irradiation, intravenous laser and laser acupuncture) were done every other day. Corticosteroid was discontinued because of its inhibitory effect on laser therapy, but the patient continued using hydroxychloroquine. After five sessions of treatment, the beneficial effects started to appear as reduction in itching and disappearing of erythema, and after 10 sessions, itching of the lesions discontinued. During the one-month follow up, the patient was symptom free and medications were discontinued. Three months later, the patient had itching at the same affected area. The LLLT with the same protocol was done for 12 sessions (phase 2).

3. Discussion

Through a total of 27 sessions of treatment, the lesions improved completely with highly satisfactory results; and one month after LLLT, as the patient was symptom free, drug-therapy was continued. Alopecia resolved completely and the atrophic lesion improved moderately (Figure 1B). After three months, the lesions flared up again with itching and scaling. A second therapeutic course was done using the same protocol for 12 sessions. During

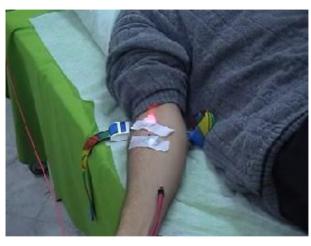


Figure 2. Intravenous Laser Therapy

the one-year follow up after the second treatment course, there was no recurrence without any medication.

Discoid lupus erythematous is a chronic inflammatory skin condition notorious for causing disfiguring scars and alopecia. Since 1986, lasers have been introduced as an alternative treatment approach for refractory skin lesions of DLE (6-8). However, lasers already been used for treatment of these lesions have been high power lasers with varying amount of adverse effects (9) (Table 1).

The primary concept for use of lasers in treating these lesions was the coagulative ability of high power lasers for blood vessels ablation and vaporization of lesions in the affected areas (6, 10, 11). However, the fact has been neglected that these lesions are sensitive to high power light irradiations such as long-standing sun exposure, UVA or UVB irradiations (7) and high power laser irradiation can also proceed the activity of disease along with other side effects such as hyper and hypo pigmentation of the lesions because of exceedingly coagulated vessels in the region and the resulting superficial ischemia.

Rather, it seems that photosensitivity of these lesions depends on various factors especially presence of some autoantibodies such as anti Ro/SS-A autoantibody (12). However, none of the previous reports have emphasized the presence of such antibodies before starting the treatment with laser irradiation.

Presence of discoid skin lesions in the process of lupus seems to be an immunological process with monouclear cell infiltrate adjacent to epidermal, adnexal, and micro vascular basement membranes and immunoglobulins and complement deposits at the basement membrane zone in about 90% of cases (3).

One of the most important functional aspects of laser

Year	Authors	Indications	Type of Laser	Response	Reported Side-Effects
1986	Henderson et al.	DLE	CO ₂	Dramatic clinical and cosmetic improvement	Splotchy hypo pigmentation
1988	Zachariae et al.	DLE	Argon	Highly satisfactory	Slight scarring and an insignificant pigmentation
1996	Nurnberg et al.	DLE	argon	Dramatic impact in contrast to untreated lesions	Hypo pigmentation
2000	Kuhn et al.	DLE	argon	Complete resolve	No short-term side-effects
2001	Tremblay JF, Carey W	DLE	Erbium: YAG	Remarkable cosmetic improvement	No side-effects during two years
2003	Baniandres et al.	DLE, CLE	FDPL, LDPL	An average improvement in over 60% of patients	Transient hyper pigmentation, light scarring

Table 1. Treatment of Discoid Lupus Erythematous With Laser; Review of the Literature

Abbreviations: CLE, cutaneous lupus erythematous; DLE, discoid lupus erythematous; FDPL, Flash lamp pulsed dye laser; LDPL, long pulsed dye laser.

therapy is photobiostimulation effects of low-level lasers on various biological systems especially immunological system, which is based on the effects of low intensity lasers, often described as lasers with less than 500 mw average power (13-17).

Low-level laser irradiation is characterized by its ability to induce non-thermal, non-destructive photo biological process, which is collectively referred to as "biomodulation" (18). Biomodulation by low level laser therapy (LLLT) has become popular as a therapeutic modality for the acceleration of wound healing and the treatment of inflammation (19). It is important to differentiate between LLLT and high-intensity laser treatment, in which heat and thermal cell damage are generated. The exact mechanism by which LLLT works is yet to be understood. However, apparently there is no linear relationship between the wavelength or dosage of LLLT and the type of photochemical reaction that it causes; depending on the target cell's physiologic status, light irradiation seems to induce either stimulatory or inhibitory effects (18).

On the other hand, LLLT appears to have immunomodulatory potential. Dyson et al. (20) investigated the effect of several energy densities of LLLT on human T-lymphocyte proliferation in vitro. Mitogen-treated cell proliferation was inhibited by all the energy densities. Interestingly, non-stimulated cells were only inhibited by higher densities in the range of 10.8 - 13.2 J/cm², while lower densities actually stimulated their proliferation (1.2 - 3.6 J/cm²). In another study, Dima et al. (21) investigated the effects of low-level laser (GaAlAs diode laser; 830 nm) on serum opsonic activity. They found that LLL irradiation had no effects on serum opsonic activity, but significantly higher values were observed at the highest dose tested (60 mw for one minute) in the absence of myeloperoxidase (MPO) inhibitor NaN₃.

To the best of our knowledge, this is the first case re-

port on the use of LLLT for the treatment of DLE. Our case, although flared after three months of withholding laser therapy, showed initial improvement without serious side effects, after six years of persistent refractory lesions and re-improvement of the lesions subsequent to second phase of therapy, fell us in doubt if we could use low-level laser therapy alone in place of drug-therapy in such refractory cases of discoid lupus erythematous, successfully. Overall, after about one year of the second therapeutic phase, the patient is still well without any medication. This encourages us to propose further clinical trials to determine the efficacy of this new modality of treatment for discoid lupus erythematous.

Acknowledgments

We acknowledge the Bahsaz laser instruments Co. Ltd, for their instrumental support and Y. Jahangiri Noudeh for the English translation of the manuscript.

References

- 1. Eastham ABW, Vleugels RA, Callen JP. Discoid lupus erythematosus 2013. Available from: http://www.emedicine.com/derm/topic247. htm.
- 2. Rudnicka L, Olszewska M, Rakowska A. Discoid Lupus Erythematosus. Atlas of Trichoscopy. Springer; 2012.
- Schmidt E, Zillikens D. Autoimmune and inherited subepidermal blistering diseases: advances in the clinic and the laboratory. *Adv Dermatol.* 1999;16:113-57. discussion 158.
- 4. Force T. Guidelines of care for cutaneous lupus erythematosus. *J Am Acad Dermatol.* 1996;**34**(5):830–6.
- Callen JP. Update on the management of cutaneous lupus erythematosus. Br J Dermatol. 2004;151(4):731-6. doi: 10.1111/j.1365-2133.2004.06196.x. [PubMed: 15491411].
- Henderson DL, Odom JC. Laser treatment of discoid lupus (case report). Lasers Surg Med. 1986;6(1):12–5. [PubMed: 3959709] 44-5.
- Spicer MS, Goldberg DJ. Lasers in dermatology. J Am Acad Dermatol. 1996;34(1):1-25.

- Garden JM, Geronemus RG. Dermatologic laser surgery. J Dermatol Surg Oncol. 1990;16(2):156–68.
- Viney C, Bachelez H, Musette P, Pinquier L, Flageul B, Dubertret L. [Cutaneous lupus erythematosus following argon laser treatment]. Ann Dermatol Venereol. 2001;128(1):49–51. [PubMed: 11226902].
- Baniandres O, Boixeda P, Belmar P, Perez A. Treatment of lupus erythematosus with pulsed dye laser. *Lasers Surg Med.* 2003;**32**(4):327–30. doi: 10.1002/lsm.10169. [PubMed: 12696102].
- Kuhn A, Becker-Wegerich PM, Ruzicka T, Lehmann P. Successful treatment of discoid lupus erythematosus with argon laser. *Dermatology*. 2000;**201**(2):175–7. [PubMed: 11053927].
- Davis TL, Lyde CB, Davis BM, Sontheimer RD. Perturbation of experimental ultraviolet light-induced erythema by passive transfer of serum from subacute cutaneous lupus erythematosus patients. *J Invest Dermatol.* 1989;92(4):573-7. [PubMed: 2703725].
- Mester A, Mester E, Mester A. Open wound healing—bedsores, ulcus cruris, burns—with systemic effects of LLLT. Lasers in Medicine and Dentistry: Basic Science and Up-to-Date Clinical Application of Low-Energy-Level Laser Therapy—LLLT. Swedish Laser-Medical Society; 2000.
- 14. Passarella S, Casamassima E, Molinari S, Pastore D, Quagliariello E, Catalano IM, et al. Increase of proton electrochemical potential and ATP synthesis in rat liver mitochondria irradiated in vitro by helium-

neon laser. FEBS Lett. 1984;175(1):95-9. [PubMed: 6479342].

- Lubart R, Wollman Y, Friedmann H, Rochkind S, Laulicht I. Effects of visible and near-infrared lasers on cell cultures. *J Photochem Photobiol B*. 1992;12(3):305–10. [PubMed: 1321905].
- Grossman N, Schneid N, Reuveni H, Halevy S, Lubart R. 780 nm low power diode laser irradiation stimulates proliferation of keratinocyte cultures: involvement of reactive oxygen species. *Lasers Surg Med.* 1998;22(4):212–8. [PubMed: 9603282].
- 17. Yu W, Naim JO, Lanzafame RJ. The effects of photo-irradiation on the secretion of TGF and PDGF from fibroblasts in vitro. *Lasers Surg Med Suppl.* 1994;**6**(8).
- 18. Schindl A, Schindl M, Pernerstorfer-Schon H, Schindl L. Low-intensity laser therapy: a review. *J Investigat Med.* 2000;**48**(5):312–26.
- Schaffer M, Bonel H, Sroka R, Schaffer PM, Busch M, Sittek H, et al. Magnetic resonance imaging (MRI) controlled outcome of side effects caused by ionizing radiation, treated with 780 nm-diode laser preliminary results. *J Photochem Photobiol B.* 2000;**59**(1-3):1-8. [PubMed: 11332876].
- 20. Dyson M, Agaiby A, Ghali L. Photobiomodulation of human Tlymphocyte proliferation in vitro. *Lasers Med Sci.* 2002;**17**(4):eA22.
- Dima VF, Suzuki K, Liu Q, Koie T, Yamada M, Suzuki KJ, et al. Laser and serum opsonic activity. *Roum Arch Microbiol Immunol.* 1996;55(4):277– 83. [PubMed: 9558962].