The Treatment of Solar Lentigo using Dr. Hoon Hur’s Optimal Melanocytic Suicide-2 Parameter with a High fluence 1064nm Nd:YAG Laser without Postinflammatory Hyperpigmentation

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Abstract
A solar lentigo is a small, well-circumscribed, pigmented macule surrounded by normal-appearing skin. Histopathologic findings may show epidermal hyperplasia and increased melanin pigmentation of the basal layer. A variable number of melanocytes are present; these melanocytes may be increased in number, but they do not form nests. In the most cases, treatment is not necessary for solar lentigo. However treating the solar lentigo without postinflammatory hyperpigmentation is very difficult and treatment for solar lentigo without side effects such as postinflammatory hyperpigmentation, scars and recurrences cannot be found in any literature yet. Therefore the authors introduce the new treatment of solar lentigo using optimal melanocytic suicide-2 parameter with a high fluence 1064nm Q-switched Nd:YAG laser without side effects or recurrences.

Introduction
A solar lentigo is a flat, sharply circumscribed patch. It can be round, oval or irregular in shape. Color varies from skin-colored, brown to dark brown or black, and size varies from a few millimeters to several centimeters in diameter [1,2]. It results from long-term exposure to Ultraviolet (UV) radiation, which causes local proliferation of epidermal melanocytes and accumulation of melanin within the keratinocytes [3]. Solar lentigos or lentigines are very common, especially in people over the age of 40 years. But the treatment of solar lentigos or lentigines without side effects such as postinflammatory hyperpigmentation (PH), scars and recurrences are extremely difficult [4-6]. Therefore the authors introduce the new treatment of solar lentigo using Dr. Hoon Hur’s optimal melanocytic suicide-2(OMS-2) parameter with a high fluence 1064nm Q-switched Nd:YAG laser without side effects or recurrences.

Report of Cases
Fifty two Korean patients with solar lentigo or solar lentigines (age range: 40-72 years old, mean age: 52.6 years) participated in this study. All patients were clinically diagnosed with solar lentigo or solar lentigines (Fig.1, 3, 5, 7, 9, 12, 15).

Figure 1: A single large round brown patch on the left lateral orbital rim area (Before treatment: 2017/1/17)

Figure 2: A complete clearance of solar lentigo (After treatment with Dr. Hoon Hur’s OMS-2 Parameter: 2017/3/2)
G-CSF Administration Accelerates Cutaneous Wound Healing Accompanied With Increased Pro-Hyp Production In db/db Mice

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Discussion

A solar lentigo (plural, solar lentigines), also known as a sun-induced freckle or senile lentigo, is a brown to dark brown lesion caused by natural or artificial ultraviolet light. Solar lentigo...
may be single or multiple [1, 2]. This type of lentigo is different from a simple lentigo (lentigo simplex) because it is caused by long-term exposure to UV light. Solar lentigo is benign, but it may indicate excessive sun exposure, a risk factor for the development of skin cancer [1, 2]. Histopathologic findings of solar lentigo may show epidermal hyperplasia and increased melanin pigmentation in the basal layer. A variable number of melanocytes are present; these melanocytes may be increased in number, but they do not form nests [3]. In the most cases, treatment is not necessary for solar lentigo. However, treating the solar lentigo without PIH is very difficult [4-6]. Although the precise pathogenesis of PIH is idiopathic, there are possible several reasons of occurrence of PIH when treating solar lentigo with conventional laser therapy. Generally 515-755nm of intense pulsed light, 532nm of Q-Switched Nd:YAG laser, 694nm of ruby laser and 755nm of alexandrite laser are absorbed in much more melanin compared to 1064nm of Q-Switched Nd:YAG laser. Thus, the laser energy which destroys epidermal melanocytes injures the surrounding keratinocytes, and the damaged keratinocytes secrete interleukin-1 (IL-1). IL-1 stimulates keratinocytes to secrete endothelin-1, α-Melanocyte Stimulating Hormone (MSH), Adrenocorticotropic Hormone (ACTH) and Prostaglandin (PGE2, PGF2α). These keratinocytic injury-induced cytokines activate melanocytes and increase melanin synthesis in the melanosomes, therefore causing PIH and worsening solar lentigo [7-11]. The damaged keratinocytes also secrete the single-chain urokinase type plasminogen activator (sc-uPA). The sc-uPA converts plasminogen to plasmin, which stimulates the keratinocytes to secrete basic fibroblast growth factor (bFGF). Again bFGF activates the melanocytes and increases melanin synthesis in the melanosomes, therefore causing PIH and worsening solar lentigo [7-11].

When the conventional laser therapy causes petechiae and crusts, laser energy may injury fibroblasts, mast cells, lymphocytes, macrophages, and vascular endothelium. Then, the fibroblasts mainly secrete Stem Cell Factor (SCF) and Hepatocyte Growth Factor (HGF) which activate melanocytes and increase melanin synthesis in the melanosomes, therefore causing PIH and worsening solar lentigo [7-11]. Finally, reactive oxygen species such as nitric oxide, free radical oxygen and peroxide, generated from the damaged keratinocytes also activate melanocytes and increase melanin synthesis in the melanosomes, eventually causing PIH and worsening solar lentigo [7-11]. To avoid the side effects such as PIH, scarring and worsening solar lentigo of the conventional laser therapy, the authors devised Dr. Hoon Hur’s optimal melanocytic suicide-2 (OMS-2) parameter therapy with a high fluence 1064nm Q-switched Nd:YAG laser without side effects or recurrences.

Dr. Hoon Hur’s OMS-2 Parameter therapy with a high 1064nm Q-switched Nd:YAG laser may destroy epidermal melanocytes without keratinocyte damage, and the end products of damaged melanocytes will be removed through transepidermal elimination [10, 11]. Also the end products of damaged melanocytes drop into the upper dermis. The dispersed melanosomes and melanins are removed by dermal melanophages through the lymphatic system [10, 11].

The authors suggest that the name of this therapy is "Dr. Hoon Hur’s OMS-2 Parameter Therapy". Because we believes that it can not only destroy epidermal melanocytes with minimal epidermal damage but also accelerate apoptotic melanocytic cell death program and can improve various skin diseases without side effects such as PIH and scarring (Table 1).

The authors think that the mechanism of Dr. Hoon Hur’s OMS-2 Parameter Therapy is to destroy the epidermal melanocytes and minimize the epidermal damage without petechiae and crusts because a wavelength of 1064nm result in less absorption by epidermal melanin. Then, weekly OMS-2 Parameter Therapy destroys melanocytes completely and accelerates apoptotic melanocyte cell death, thus the end products of damaged melanocytes such as the dispersed melanosomes and melamins are either removed by the transepidermal elimination or are removed by dermal melanophages through the lymphatic system [10, 11]. Eventually, complete clearance of solar lentigo without side effects and recurrences can be achieved. In these cases, our patients with solar lentigo were treated with 8 sessions of a high fluence 1064nm Q-switched Nd:YAG laser (QX-MAX Laser, Fotona, Slovenia) at a one-week interval with a spot size of 3mm, a fluence of 5J/cm² and a pulse rate of 10Hz with pulse stacking technique for 3 seconds to the solar lentigo. The pulse stacking technique for 3 seconds to the solar lentigo is very important to destroy the epidermal melanocytes with minimizing the epidermal damage. If the pulse stacking technique for more than 3 seconds to the solar lentigo was performed, the epidermal damages might have occurred and the damaged keratinocytes might have secreted the keratinocytic injury-induced cytokines such as endothelin-1, α-MSH, ACTH, bFGF, and prostaglandin (PGE2, PGF2α) which activate melanocytes and increase melanin synthesis in the melanosomes, therefore causing PIH and worsening solar lentigo [7-11]. In case of solar lentigo on face, 8 sessions of OMS-2 Parameter Therapy are performed once a week. But in case of solar lentigo on other parts of body (arms, legs, and torso), more than 8 sessions of OMS-2 Parameter Therapy should be performed once a week regardless of the lesion size. The merit of OMS-2 Parameter Therapy is that it minimizes the epidermal damage without petechiae and crusts. This parameter can deliver the sufficient energy to destroy epidermal melanocytes and also salvage normal background tissue due to less absorption by epidermal mela-
nin. Therefore this therapy does not provoke PIH and scarring. But OMS-2 Parameter Therapy requires 8 treatment sessions for 8 weeks. In these cases, 52 patients with a solar lentigo or solar lentigines (Fig. 1,3,5,7,9,12,15) were treated with OMS-2 Parameter of a high fluence 1064nm Q-switched Nd:YAG laser. A total of 52 patients with a solar lentigo or solar lentigines was achieved complete clearance of the pigmented lesions without PIH (Fig. 2, 4, 6, 8, 10, 13, 16). There are no recurrences at 18 months follow-up (Fig.11, 14). All patients were satisfied with the results of Dr. Hoon Hur’s OMS-2 Parameter Therapy, and no any significant side effects, including PIH and scarring.

Conclusion

The parameter for each Dr. Hoon Hur’s OMS-2 Parameter Therapy was a spot size of 3mm, a fluence of 5J/cm2 and a pulse rate of 10Hz with pulse stacking technique for 3 seconds. This therapy does not cause side effects such as petechiae, crusts and PIH during the laser treatment. There is no recurrence at 18 months follow-up. Therefore, Dr. Hoon Hur’s OMS-2 Parameter Therapy is thought that more effective and safe results of treatment than conventional laser treatment can be expected.

References