ORIGINAL RESEARCH

Assessment of efficacy Monochromatic Excimer Light in the treatment of psoriasis

Dr. Rajneesh Kanwat

Associate Professor, Department of Dermatology, United Institute of Medical Sciences, Prayagraj, Uttar Pradesh, India

Corresponding Author

Dr. Rajneesh Kanwat

Associate Professor, Department of Dermatology, United Institute of Medical Sciences, Prayagraj, Uttar Pradesh, India

Received: 13 February, 2022 Accepted: 15 March, 2022

ABSTRACT

Background: Psoriasis is a chronic autoimmune skin disorder characterized by the rapid growth of skin cells, leading to the formation of thick, scaly patches. The present study was conducted to assess the clinical efficacy of Monochromatic Excimer Light (MEL) in the treatment of psoriasis. **Materials & Methods:** 74 cases of psoriasis of both genderswere treated by MEL, two sittings per week. PSI was calculated before and after treatment which was derived from the standard Psoriasis Area and Severity Index (PASI) by omitting the area, thus assigning a score of 0-4 (0=none; 1=mild; 2=significant; 3=moderate; 4=severe) for erythema, induration and desquamation. **Results:** Out of 74 patients, 30 were males and 44 were female. Common site was scalp seen in 35, trunk in 38, palms in 12, upper limb in 67 and lower limb in 71 cases. Outcome was excellent in 62, good in 10 and moderate in 2 cases. The difference was significant (P< 0.05). **before** treatment PASI score was 7.4 and after treatment was 1.3. The difference was significant (P< 0.05). **Conclusion:** MEL is a safe and efficient treatment for plaque psoriasis, and the results are primarily dependent on the patient's skin phototype and the anatomic location of the lesion.

Keywords: Autoimmune, Psoriasis, skin disorder

This is an open access journal, and articles are distributed under the terms of the Creative Commons Attribution- Non Commercial-Share Alike 4.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as appropriate credit is given and the new creations are licensed under the identical terms.

INTRODUCTION

Psoriasis is a chronic autoimmune skin disorder characterized by the rapid growth of skin cells, leading to the formation of thick, scaly patches. It is a long-lasting condition that can vary in severity, from small localized patches to complete body coverage. Psoriasis is not contagious, and its exact cause is still not fully understood, though it involves an interplay of genetic, environmental, and immune system factors. 3

Plaque Psoriasis (Psoriasis Vulgaris) is the most common form, accounting for about 80-90% of cases. Symptoms are raised, red patches covered with a silvery-white buildup of dead skin cells (scales). These patches, or plaques, are often itchy or sore and typically appear on the elbows, knees, scalp, and lower back. Guttate Psoriasis often occurs in children and young adults, sometimes after a bacterial infection like strep throat. Symptoms are small, drop-shaped lesions on the trunk, arms, and legs. These spots are usually not as thick as plaque psoriasis. 5

For mild disease, topical medications are usually utilized, phototherapy for moderate disease, and systemic medications for severe disease. A kind of

ultra violet lamp known as a monochromatic excimer light (MEL) uses argon, krypton, xenon, and halogen among other noble gases.⁶ High pressure combined with the right electrical stimulation results in the pseudo-molecule MEL, which has UV laser light activity. MEL is a new type of UV-B light used in dermatology to treat conditions like vitiligo, alopecia areata, psoriasis, atopic dermatitis, and mycosis fungoides in the patch stage.⁷The present study was conducted to assessthe clinical efficacy of Monochromatic Excimer Light (MEL) in the treatment of psoriasis.

MATERIALS & METHODS

The present study was conducted on 74 cases of psoriasis of both genders. All gave their written consent to participate in the study.

Data such as name, age, gender etc. was recorded. Psoriasis history, medical history, treatments received for psoriasis and other treatments were recorded for each patient. All patients were treated by MEL, two sittings per week. PSI was calculated before and after treatment which was derived from the standard Psoriasis Area and Severity Index (PASI) by omitting

the area, thus assigning a score of 0-4 (0=none; 1=mild; 2=significant; 3=moderate; 4=severe) for erythema, induration and desquamation. Results thus

obtained were subjected to statistical analysis. P value < 0.05 was considered significant.

RESULTS

Table I Distribution of patients

Total- 74				
Gender	Male	Female		
Number	30	44		

Table I shows that out of 74 patients, 30 were males and 44 were female.

Table II Assessment of parameters

Parameters	Variables	Number	P value
Site	Scalp	35	0.85
	Trunk	38	
	Palms	12	
	Upper limb	67	
	Lower limb	71	
Outcome	Excellent	62	0.01
	Good	10	
	Moderate	2	
	Poor	0	

Table II, graph I shows that common site was scalp seen in 35, trunk in 38, palms in 12, upper limb in 67 and lower limb in 71 cases. Outcome was excellent in 62, good in 10 and moderate in 2 cases. The difference was significant (P< 0.05).

Graph I Assessment of parameters

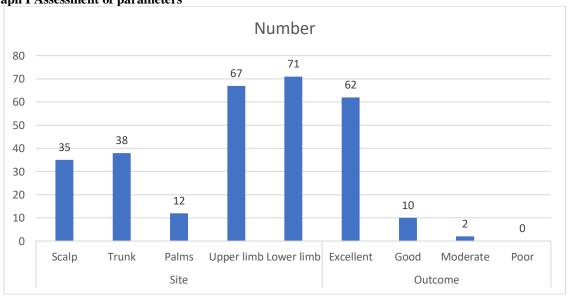


Table III Assessment of PASI

PASI	Mean	P value
Before treatment	7.4	0.01
After treatment	1.3	

Table III shows that before treatment PASI score was 7.4 and after treatment was 1.3. The difference was significant (P < 0.05).

DISCUSSION

Psoriasis is believed to result from an overactive immune system that causes inflammation and accelerated skin cell turnover.^{8,9} This leads to the buildup of skin cells on the surface, forming the characteristic scales and plaques.^{10,11}A family history

of psoriasis increases the risk, infections such as strep throat or skin infections, stress can trigger or exacerbate symptoms, cold weather tends to worsen the condition, injury to the skin such as cuts, scrapes, or sunburn (Koebner phenomenon), certain drugs, like lithium or beta-blockers, can trigger psoriasis, alcohol

and smokingcan exacerbate the condition, hormonal changes particularly in women during puberty, pregnancy, or menopause. 12,13 The present study was conducted to assess the clinical efficacy of Monochromatic Excimer Light (MEL) in the treatment of psoriasis.

We found that out of 74 patients, 30 were males and 44 were female. Mohamed et al14 evaluated the clinical efficacy and safety of Monochromatic Excimer Light (MEL) in the treatment of psoriasis. Ninety Egyptian psoriasis patients with plaque-type psoriasis were treated with MEL twice a week, for a total of 24 sessions or until complete improvement. The Psoriasis Severity Index (PSI) was assessed for each patient. At the final visit, 96/360 (26.67%) patches of psoriasis showed complete clearance, 75/360 (20.83%) patches achieved more than 75% improvement, 132/360 (36.67%) patches achieved 51-75% improvement and (15.83%) patches achieved 26-50% improvement. The best result occurred on the trunk followed by lower limbs, upper limbs, palms and soles than the scalp. There was inverse statistical correlation between the percentage of improvement and skin phototype and no correlation with patient's age, sex or duration of disease. Side effects reported were erythema in 45 patches, post inflammatory hyperpigmentation in 102 patches, perilesional skin was covered by protective sheet, so no side effect occurred. No recurrence was observed six months after treatment.

We found that common site was scalp seen in 35, trunk in 38, palms in 12, upper limb in 67 and lower limb in 71 cases. Outcome was excellent in 62, good in 10 and moderate in 2 cases. Goldinger SM et al¹⁵determined the response of psoriasis lesions to the 308-nm excimer laser compared to 311-nm UVB phototherapy. They selected psoriasis plaques of 16 patients were treated with the excimer laser whereas the rest of the body was treated with UVB narrowband phototherapy. A modified PASI score was used to evaluate the results. After 12 treatments, 15 patients were evaluated. In 2 patients no difference between the two body sides was observed. In 9 patients the laser-treated lesions showed better results, whereas in 4 patients the side treated with 311-nm UVB showed more clearing. The mean reduction in PASI score was 5.6 and 4.9, respectively (difference significant). The use of the 308-nm xenon chloride excimer laser is an additional effective therapeutic option for the treatment of psoriasis vulgaris.

We found that before treatment PASI score was 7.4 and after treatment was 1.3. Kollner K et al¹⁶enrolled fifteen patients with plaque psoriasis. Three different psoriatic lesions were treated with the 308-nm excimer laser, the 308-nm excimer lamp or 311-nm narrowband UVB three times per week. Sixteen patients were enrolled in the second regime. Two plaques were treated with the 308-nm excimer laser or with the 308-nm lamp with an accelerated scheme three times per week. Using Friedman analysis, the

PSI scores did not show a statistically significant difference (P>0.05) comparing 308-nm laser therapy, 308-nm lamp therapy and 311-nm narrowband therapy after 10 weeks in the first regime. The mean number of treatments to achieve clearance was 24. With the accelerated scheme, clearance could be achieved with fewer treatments and with half the cumulative dose of the first regime. Nevertheless, the side-effects such as blistering and crusting were also increased.

The shortcoming of the study is small sample size.

CONCLUSION

Authors found that MEL is a safe and efficient treatment for plaque psoriasis, and the results are primarily dependent on the patient's skin phototype and the anatomic location of the lesion.

REFERENCES

- Menter A, Griffiths CE. Current and future management of psoriasis. Lancet. 2007;370(9583):272-84.
- Kemeny L, Bonis B, Dobozy A, Bor Z, Szabo G, Ignacz F, et al. 308-nm excimer laser therapy for psoriasis. Arch Dermatol. 2001;137:95-96.
- Asawanonda P, Anderson RR, Chang Y, Taylor CR. 308-nm excimer laser for the treatment of psoriasis: a dose-response study. Arch Dermatol. 2000;136:619-24.
- Van Weelden H, Baart de la Faille H, Young E, van der Leun JC. Comparison of narrow-band UV-B phototherapy and PUVA photo-chemotherapy in the treatment of psoriasis. Acta Derm Venereol. 1990;70:212-15.
- Trehan M, Taylor CR. High-dose 308 nm excimer laser for the treatment of psoriasis. J Am Acad Dermatol. 2002;46:732-37.
- 6. Rodewald EJ, Housman TS, Mellen BG, Feldman SR. Follow-up survey of 308- nm laser treatment of psoriasis. Lasers Surg Med. 2002;31:202-06.
- Feldman SR, Mellen BG, Housman TS, Fitzpatrick RE, Geronemus RG, Friedman PM, et al. Efficacy of the 308 nm excimer laser for the treatment of psoriasis: results of a multicenter study. J Am Acad Dermatol. 2002;46:900-06.
- 8. Spann CT, Barbagallo J, Weinberg JM. A review of the excimer laser in the treatment of psoriasis. Cutis. 2001;68:351-52.
- 9. Spencer JM, Hadi SM. The excimer lasers. J Drugs Dermatol. 2004;3:522-25.
- Holroyd RA, Cook AR, Preses JM. Dynamics of excimer formation and decay in supercritical krypton. J Chem Phys. 2009;131(22):224509.
- Kemeny L, Bonis B, Dobozy A, Bor Z, Szabo G, Ignacz F, et al. 308-nm excimer laser therapy for psoriasis. Arch Dermatol. 2001;137:95-96.
- 12. Mavilia L, Mori M, Rossi R, Campolmi P, Puglisi Guerra A, Lotti T, et al. 308 nm monochromatic excimer light in dermatology: personal experience and review of the literature. G Ital Dermatol Venereol. 2008;143:329-37.
- Wolff K, Gschnait F, Honigsmann H, Konrad K, Parrish JA, Fitzpatrick TB, et al Phototesting and dosimetry for photochemotherapy. Br J Dermatol. 1977;96:1-10

- 14. Mohamed EM, Tawfik KM, Mohamed RR, Abdelgwad AM, Elsaie ML. Treatment of Plaque Psoriasis by 308 nm Excimer Light in Egyptian Patients. Journal of clinical and diagnostic research. 2018 Mar 1;12(3):01-4.
- Goldinger SM, Dummer R, Schmid P, Prinz Vavricka M, Burg G, Lauchli S, et al. Excimer laser versus narrow-band UVB (311 nm) in the treatment of psoriasis vulgaris. Dermatology. 2006;213(2):134-39.
- Kollner K, Wimmershoff MB, Hintz C, Landthaler M, Hohenleutner U. Comparison of the 308-nm excimer laser and a 308-nm excimer lamp with 311-nm narrowband ultraviolet B in the treatment of psoriasis. Br J Dermatol. 2005;152:750-54.