ORIGINAL ARTICLE

Comparison of efficacy & safety of Blue Light vs Topical Application of 1% Clindamycin solution in the treatment of mild to moderate inflammatory Acne Vulgaris

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ABSTRACT

Aim: To compare the efficacy & safety of blue light vs topical application of 1% clindamycin solution in the treatment of mild to moderate inflammatory acne vulgaris.

Study Design: Comparative interventional study done in Dermatology Outpatient Department Unit-II, KEMU/ Mayo Hospital, Lahore six months i.e. 1-06-2014 – 30-11-2014

Methodology: After an informed and written consent, 130 patients fulfilling the selection criteria were enrolled in the study and divided in two study groups A & B by balloting method. At first visit, a detailed history and clinical examination was recorded on a specially designed proforma. The acne was graded according to the acne grading scale of American Academy of Dermatology.^{4,13} Group A was exposed to blue light for twenty minutes twice weekly for eight weeks. Group B was given 1% clindamycin to apply twice daily for a period of eight weeks. Post- treatment follow up was done for next four weeks. Patients were assessed at 2nd, 4th, 6th, 8th, 10th, and 12th week. All findings and side effects were recorded on a predesigned proforma. To determine the efficacy of treatment, Acne Severity Index (ASI) was used.⁵

Result: Efficacy of treatment [≥ 50% improvement in ASI score] was seen in 39(60%) patients in blue light group while in clindamycin group it was achieved in 8 (12.3%) patients only. Blue light group had significantly less number of side effects observed in 35 (37.23%) patients while clindamycin group had a higher number of side effects observed in 59 (62.77%) patients, p-value= 0.013.

Conclusion: Blue light is more efficacious and safer than topical 1% clindamycin in the treatment of mild to moderate inflammatory acne vulgaris.

Keywords: Blue light, 1% Clindamycin, Acne vulgaris

INTRODUCTION

Acne vulgaris is a common, chronic inflammatory disease of pilosebaceous unit affecting 80% of teenagers between 13-18 years and 50.9% of women and 42.5% of men between the ages of 20 to 29 years 1,2,3. Topical and systemic antibiotics are mainstay of treatment but there is a rapid increase in the resistance to antibiotics for Propionibacterium acnes, so, there is a need for some alternative therapy.5Phototherapy with blue light is an efficacious & safe additional therapy⁶. Blue light treatment is a non-UV light therapy, ranging between the wavelength of 405-485nm.7 It is a natural, non-invasive treatment and effective against those strains of Propionibacterium acnes which are resistant to antibiotics⁷. There is photo-excitation of bacterial porphyrins after exposure to blue visible light, due to which singlet oxygen radical is produced which endogenous photodynamic destruction bacteria^{5,6}. Light is absorbed by bacterial cells which produces changes in permeability of cell membrane. This change in permeability leads to influx of proton and disruption of pH gradient across cell membrane which leads to inhibition of growth of P. Acnes. 5,6 Inhibition of

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growth and photodynamic demolition of *P. acnes*, both play an important role in the treatment of inflamed acne lesions by blue light. 6Clindamycin is a lincosamide antibiotic, active against many gram-positive cocci, anaerobic and microaerophilic gram-positive & gram-negative organisms including Propionibacteriumacnes.8 Its bactericidal action is due to its capability to inhibit ribosomal protein synthesis in target organisms by binding to 50S ribosomal subunits.8 Different studies have shown excellent results ranging from 34% to 76% betterment in acne lesions with blue light therapy^{5,6,9-12}. Morton CA et al did a study which showed the average clearance of acne lesions upto 76% with blue light after 4 weeks of treatment. 6 Shalita AR et al studied the effect of blue light in the treatment of mild to moderate acne vulgaris.9 After eight bi-weekly treatments, inflammatory lesion count decreased upto 60%. Two weeks after last treatment, count further decreased to almost 70%9. Gold MH et al compared the efficacy of blue light with topical 1% clindamycin in the treatment of mild to moderate inflammatory acne vulgaris.11 Blue light therapy decreased inflammatory lesions count by an average of 34%, as compared to 14% for 1% clindamycin solution¹¹.

This study was planned to improve the management and provide the patient with best effective & safe treatment for acne.

METHODOLOGY

After approval from ethical committee of KEMU, the study was conducted in Dermatology Outpatient Department Unit-II, KEMU/ Mayo Hospital, Lahore from i.e. 1-06-2014 -30-11-2014. After an informed and written consent, 130 patients fulfilling the selection criteria were enrolled in the study and divided in two study groups A & B by balloting method. At first visit, a detailed history and clinical examination was recorded on a specially designed proforma. The acne was graded according to the Acne Grading Scale of American Academy of Dermatology. 4,13 Lesions were counted before starting therapy and photographs were taken at each follow-up visit during & after treatment. To determine the efficacy of treatment, Acne Severity Index (ASI) was used.⁵ Patients were instructed to cleanse their face before each treatment with an unscented soap or facial cleanser. They were instructed to apply a moisturizing non comedogenic sunscreen with sun protection factor (SPF) 30 after each morning treatment. Group A was exposed to blue light for twenty minutes two times per week (3-4 days interval between treatments) for eight weeks. Blue light source was kept at 5-10 cm from patient's face and eyes of the patients were covered with black goggles. Group B was given 1% clindamycin solution to apply twice daily for a period of eight weeks. After eight weeks, follow up was done for next four weeks. Patients were assessed at 2nd, 4th, 6th, 8th, 10th and 12th week in order to compare the efficacy and safety Acne Grading Scale

of blue light with 1% clindamycin antibiotic solution. All findings and side effects were recorded on a predesigned proforma.

ASI score= 0.25 x comedone number + 1 x papule number + 2 x pustule number

Assessment criteria: Criteria of improvement was as follows:

- ullet <25% reduction in ASI score was regarded as poor improvement
- 25-49% reduction in ASI score was regarded as fair improvement
- 50-75% reduction in ASI score was regarded as good improvement
- >75% reduction in ASI score was regarded as excellent improvement

Data Analysis Procedure: Data entry and analysis was done by using SPSS 17. Quantitative variables like age, number of acne lesions, duration of disease etc., was presented by using mean and standard deviation (SD). Qualitative variables like sex of patient, types of acne lesions etc., were presented by using frequency tables, percentages and appropriate graphs. Repeated measure ANOVA / Friedman test was used to see the reduction in ASI score from the baseline till last follow-up. Clinical improvement and adverse effects of both treatments were compared by using chi-square test. A p-value of ≤0.05 was taken as significant.

Acne	Comedones	Papules	Pustules	Nodules/ Cysts	Scar
Mild	Several-many	Several	Few	None	No
Moderate	Several-numerous	Several	Several	Rare	No
Severe	Numerous/Scattered	Numerous	Numerous	Many	Often

RESULTS

Table 1: Comparison of age (years) in both study groups

		Mean S.D.		95% C.I. for Mean	
		Wieaii	3.D.	Lower	Upper
	Blue light (n=65)	21.89	4.79	20.71	23.08
Age (years)	Clindamycin(n=65)	23.68	5.92	22.21	25.14
	Total(n=130)	22.78	5.44	21.84	23.73

Table 2: Comparison of duration of disease (years) in both study groups

		Mean	S.D.	95% C.I. for Mean		
		Weari	3.D.	Lower	Upper	
Duration of disease (years)	Blue light (n=65)	2.14	2.23	1.59	2.70	
	Clindamycin(n=65)	1.63	1.65	1.22	2.04	
	Total(n=130)	1.89	1.97	1.55	2.23	

Table 3: Comparison of comedones in both study groups

		Mass	Maan CD	Mean S.D	95% C.I. for M	ean
		wean	3.0	Lower	Upper	
	Blue light	26.86	14.76	23.20	30.52	
No. of lesions (before treatment)	Clindamycin	14.06	7.61	12.18	15.95	
	Total	20.46	13.35	18.15	22.78	
	Blue light	26.74	14.44	23.16	30.32	
No. of lesions 2nd week	Clindamycin	14.14	8.07	12.14	16.14	
	Total	20.44	13.26	18.14	22.74	
No. of lesions 4th week	Blue light	25.97	13.23	22.69	29.25	
	Clindamycin	14.57	8.06	12.57	16.57	
	Total	20.27	12.32	18.13	22.41	

No of losions	Blue light	25.95	13.46	22.62	29.29
No. of lesions 6th week	Clindamycin	14.42	7.92	12.45	16.38
our week	Total	20.18	12.43	18.03	22.34
	Blue light	25.69	13.58	22.33	29.06
No. of lesions 8th week	Clindamycin	14.31	7.90	12.35	16.26
	Total	20.00	12.45	17.84	22.16
	Blue light	25.89	12.94	22.69	29.10
No. of lesions 10th week	Clindamycin	14.09	7.81	12.16	16.03
	Total	19.99	12.18	17.88	22.11
	Blue light	27.08	13.67	23.69	30.46
No. of lesions12th week	Clindamycin	13.77	9.00	11.54	16.00
	Total	20.42	13.32	18.11	22.73
	Blue light	7.03	29.18	0.38	14.44
% reduction in no. of lesions	Clindamycin	8.09	54.30	6.06	22.24
	Total	7.54	43.11	0.21	15.30

Table 4: Comparison of papules in both study groups

		Mean	6.5	95% C.I. fo	or Mean	
			S.D	Lower	Upper	p-value
	Blue light (n=65)	13.75	12.55	10.65	16.86	
No. of lesions (before treatment)	Clindamycin(n=65)	9.40	3.45	8.55	10.25	0.008
	Total(n=130)	11.58	9.42	9.94	13.21	
	Blue light (n=65)	11.20	4.00	10.21	12.19	
No. of lesions 2nd week	Clindamycin(n=65)	8.77	2.74	8.09	9.45	< 0.001
	Total(n=130)	9.98	3.63	9.36	10.61	
	Blue light (n=65)	9.17	3.37	8.33	10.00	
No. of lesions 4th week	Clindamycin(n=65)	7.78	2.32	7.21	8.36	0.007
	Total(n=130)	8.48	2.97	7.96	8.99	
	Blue light (n=65)	7.62	3.18	6.83	8.40	0.111
No. of lesions 6th week	Clindamycin(n=65)	6.86	2.07	6.35	7.37	
	Total(n=130)	7.24	2.70	6.77	7.71	
	Blue light (n=65)	5.15	2.35	4.57	5.74	
No. of lesions 8th week	Clindamycin(n=65)	6.51	3.26	5.70	7.31	0.007
	Total(n=130)	5.83	2.91	5.33	6.34	
	Blue light (n=65)	4.80	2.43	4.20	5.40	
No. of lesions 10th week	Clindamycin(n=65)	6.82	2.14	6.28	7.35	<0.001
	Total(n=130)	5.81	2.50	5.37	6.24	
	Blue light (n=65)	4.85	2.24	4.29	5.40	
No. of lesions 12th week	Clindamycin(n=65)	6.69	2.11	6.17	7.21	0.001
	Total(n=130)	5.77	2.35	5.36	6.18	
	Blue light (n=65)	61.68	9.78	59.25	64.10	
% reduction in no. of lesions	Clindamycin(n=65)	27.28	16.94	23.09	31.48	<0.0001
	Total(n=130)	44.48	22.08	40.65	48.31	

Table 5: Comparison of pustules in both study groups

		Mean	S.D	95% C.I. fo	or Mean	n volue
		wean	ა.ს	Lower	Upper	p-value
	Blue light (n=65)	6.43	3.48	5.57	7.29	
No. of lesions (before treatment)	Clindamycin(n=65)	7.20	2.68	6.54	7.86	0.161
(Total(n=130)	6.82	3.12	6.27	7.36	
	Blue light (n=65)	6.06	2.94	5.33	6.79	
No. of lesions 2nd week	Clindamycin(n=65)	7.37	3.25	6.56	8.18	0.018
	Total(n=130)	6.72	3.16	6.17	7.26	0.0.0
	Blue light (n=65)	4.85	2.84	4.14	5.55	
No. of lesions 4th week	Clindamycin(n=65)	6.29	3.31	5.47	7.11	0.009
	Total(n=130)	5.57	3.16	5.02	6.12	
	Blue light (n=65)	3.91	2.75	3.23	4.59	
No. of lesions 6th week	Clindamycin(n=65)	5.65	4.23	4.60	6.69	0.006
	Total(n=130)	4.78	3.66	4.14	5.41	

	Blue light (n=65)	2.22	2.29	1.65	2.78	
No. of lesions 8th week	Clindamycin(n=65)	4.28	3.32	3.45	5.10	< 0.001
	Total	3.25	3.03	2.72	3.77	
	Blue light	1.65	1.80	1.20	2.09	
No. of lesions 10th week	Clindamycin	4.23	2.64	3.58	4.89	< 0.001
	Total	2.94	2.60	2.49	3.39	
	Blue light	2.32	2.11	1.80	2.85	
No. of lesions 12th week	Clindamycin	4.38	1.68	3.97	4.80	< 0.001
	Total	3.35	2.16	2.98	3.73	
% reduction in no. of lesions	Blue light	70.09	26.10	63.63	76.56	
	Clindamycin	37.86	13.43	34.54	41.19	<0.0001
	Total	70.09	26.10	49.42	58.53	

Table 6: Comparison of asi scores in both study groups

		Mean	S.D.	95% C.I. f	95% C.I. for Mean	
		Iviean	S.D.	Lower	Upper	p-value
	Blue light (n=65)	31.01	12.13	28.00	34.02	
ASI score before treatment	Clindamycin(n=65)	26.58	7.03	24.84	28.32	0.012
	Total(n=130)	28.80	10.13	27.04	30.55	
	Blue light (n=65)	30.08	12.14	27.07	33.09	
ASI score 2nd week	Clindamycin(n=65)	26.07	7.11	24.31	27.84	0.023
	Total(n=130)	28.08	10.11	26.32	29.83	
	Blue light (n=65)	25.40	11.03	22.67	28.13	
ASI score 4th week	Clindamycin(n=65)	22.96	6.42	21.37	24.55	0.125
	Total(n=130)	24.18	9.07	22.60	25.75	
	Blue light (n=65)	21.88	10.66	19.24	24.52	0.171
ASI score 6th week	Clindamycin(n=65)	19.83	5.59	18.44	21.21	
	Total(n=130)	20.85	8.54	19.37	22.34	
	Blue light (n=65)	15.67	8.21	13.64	17.71	
ASI score 8th week	Clindamycin(n=65)	17.18	4.88	15.97	18.39	0.206
	Total(n=130)	16.43	6.77	15.25	17.60	
	Blue light (n=65)	14.60	7.56	12.73	16.48	
ASI score 10th week	Clindamycin(n=65)	18.28	5.08	17.02	19.54	0.001
	Total(n=130)	16.44	6.67	15.28	17.60	
	Blue light (n=65)	15.25	7.70	13.24	17.26	
ASI score 12th week	Clindamycin(n=65)	19.24	5.76	17.81	20.67	0.001
	Total(n=130)	17.34	7.02	16.09	18.59	
	Blue light (n=65)	51.55	7.64	49.66	53.45	
ASI reduction(%)	Clindamycin(n=65)	26.32	5.17	25.04	27.60	< 0.001
	Total(n=130)	38.94	14.23	36.47	41.41	

Table 7: Comparison of efficacy in both study groups

		Stud	Total	
		Blue light	Clindamycin	lotai
V		39	8	47
Yes	res	60.0%	12.3%	36.2%
Efficacy	No	26	57	83
	No	40.0%	87.7%	63.8%
T-1-1		65	65	130
Total		100.0%	100.0%	100.0%

p-value< 0.001

Table 8: Comparison of side effects in both study groups

	Stu	Study groups		
	Blue light Clindamycin		p-value	
Burning 2nd week	0	1	0.315 (insignificant)	
Burning 4th week	0	3	0.080 (insignificant)	
Burning 6th week	3	4	0.689 (insignificant)	
Burning 8th week	0	7	0.007 (significant)	
Burning 8 th -12 th week	0	0		

Itching 2nd week	4	4	1 (insignificant)
Itching 4th week	13	21	0.11 (insignificant)
Itching 6th week	23	34	0.052 (insignificant)
Itching 8th week	10	29	0.000 (significant)
Itching 10th week	0	11	0.001 (significant)
Itching 12th week	0	3	0.80 (insignificant)
Dryness 2nd week	1	0	0.315 (insignificant)
Dryness 4th week	17	8	0.045 (significant)
Dryness 6th week	26	31	0.377 (insignificant)
Dryness 8th week	10	35	<0.001 (significant)
Dryness 10th week	0	3	0.080 (insignificant)
Dryness 12th week	0	2	0.154 (insignificant)
Erythema 4th week	6	3	0.30 (insignificant)
Erythema 6th week	4	15	0.006 (significant)
Erythema 8th week	3	28	<0.001 (significant)
Erythema 10th week	0	1	0.135 (insignificant)
Peeling 6th week	0	3	0.80 (insignificant)
Peeling 8th week	0	1	0.315 (insignificant)
Any Other 6th week	0	2	0.154 (insignificant)

DISCUSSION

This study was planned to improve the management and provide the patient with best effective & safe treatment for acne. The mean age of patients was 22.78 ± 5.44 years in our study, which correlates with the study by Wheeland RG in which mean age of patients was 22 ± 6.7 years. ¹⁰ These results show that acne is more prevalent among this age group. This also correlates well with the study performed by Collier CN *et al* which shows that acne affects at higher rates between 20-29 years. ³ Acne Grading Scale of American Academy of Dermatology was used in this study to classify patients having mild to moderate inflammatory acne. Same grading scale was used by Rahman MM *et al* in his study¹³.

The results of our study show that mean number of comedonal lesions was statistically same from baseline till 12th week in both groups. No significant difference was found in % reduction of comedonal lesions between both groups. This is in accordance with the study performed by Morton CA *et al* which showed that Blue light treatment had little effect on the number of comedones. ⁶It also correlates well with the study performed by Lookingbill DP *et al* which showed that topical Clindamycin alone is not effective in reducing non-inflammatory lesions ¹⁴.

The results of our study show that in Blue light group the mean % reduction in the number of papules and pustules was 61.68 and 70.09 respectively. This result is in accordance with the studies performed by Shalita and Papageorgiou which showed that Blue light is effective in reducing inflammatory lesions of acne vulgaris (60%)^{15,16}.

Morton CA observed the average reduction in inflammatory lesions was upto 73% with Blue light.⁶ Wheeland RG showed that Blue light is effective upto 46% in reducing the inflammatory lesions.¹⁰ Similarly Gold MH showed that Blue light is effective upto 41.03% in reducing the inflammatory lesions¹². This difference in the results of various studies could be explained by the fact that different

study tools were used in these studies and there may be different response pattern among study groups and the time to reach the optimum clearance differed between subjects. The results of our study show that in Clindamycin group mean % reduction in the number of papules and pustules was 27.28 was 37.86 respectively. This result is in accordance with the study performed by Lookingbill DP *et al* which showed that reduction of inflammatory lesions was upto 35% with Clindamycin¹⁴.

Present study reported that efficacy of Blue light was significantly higher when compared to Clindamycin in terms of reduction of ASI score as (51.55% vs. 26.32%). Similar study was performed by Gold MH *et al* which showed that Blue light was more effective than topical 1% Clindamycin as (34% vs. 14%)¹¹.

Present study showed that Blue lightis moresafe than Clindamycin. As 37% patients experienced side effects in Blue light group and 62% patients in Clindamycin group. Most frequently observed side effects were itching and dryness. Mild erythema was observed in few patients. Arruda LHF observed that only 23.3% patients hadmild adverse effects characterized by desquamation and dryness with Blue light treatment⁹. Wheeland RG also show that53 percent of subjects agreed that the Blue light treatment has less side effects than traditional acne treatments. Most frequently observed side effect was minimal and transient skin dryness¹⁰. Faghihi G et al observed dryness (18%) and mild erythema (25%) with Blue light treatment.⁵

So according to the result of our study we propose that Blue light therapy significantly reduces inflamed acne lesions, with mild side effects, and offers an effective and safe treatment for acne vulgaris. This study is consistent with previous reports of Blue light used in acne and suggests that Blue light phototherapy deserves inclusion in the list of therapeutic options for patients with mild to moderate acne. Therapies that avoid oral ingestion of medication and minimize topical applications are likely to

be popular with patients. But compliance is major problem because patient has to go to the treatment center twice a week to undergo Blue light applications. So further studies are required to observe optimum efficacy with the shortest possible duration and number of treatments. We also suggest that further studies are required as regards the use of other treatment options in conjunction with Blue light therapy to provide an effective therapy combination.

CONCLUSION

Blue light therapy is effective, safe and favorable as compared to topical 1% clindamycin in patients with mild to moderate acne. Its gentleness on the skin offers a better choice in patients who cannot use antibiotics or topical irritating therapies or it may be used as an adjuvant therapy.

Conflict of interest: Nil

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