M89: A COMBINATION OF 89% MINERALIZED THERMAL WATER AND HYALURONIC ACID IS EFFECTIVE AND WELL TOLERATED AFTER DERMATOLOGIC PROCEDURES

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Table 1

INTRODUCTION

The skin exposome comprises several external and internal factors including UV radiation, climatic conditions (heat and humidity), medications, pollution, stress, and dermatologic procedures that may damage the skin barrier, induce skin diseases or accelerate skin ageing.1-5

Mineral 89 (M89, Vichy Laboratoires), containing 89% Vichy volcanic water and hyaluronic acid in a minimalist formulation, was developed to reinforce the skin barrier and to protect against exposome factors.6-9

Recent study results from 1630 subjects with inflammatory dermatoses or having undergone dermatologic procedures confirmed the benefit and excellent tolerance of M89.10

AIM OF THE STUDY

The aim of this study was to assess after 4 weeks of daily use the efficacy and tolerability of M89 in adult subjects having undergone dermatologic procedures.

METHODOLOGY

In an international, multicenter observational study, subjects having undergone dermatologic procedures applied for 4 weeks once or twice daily M89. Data about dermatologic procedures (including acne, scars, skin ageing, spots), subject information, skin characteristics compliance, subject perception of efficacy, tolerance, as well as investigator satisfaction were collected after 4 weeks. Subject satisfaction was assessed after 1 and 4 weeks of use.

Table 2

RESULTS

Data from 1101 subjects were analyzed. Peeling accounted for 23.3% and Laser/IPL for 22.1% of dermatologic procedures. At baseline, subjects had mainly dry (47.6%) and sensitive skin (60.5%). 50.3% presented with some degree of erythema, 54.0% with desquamation and 63.8% with irritation. A total of 56.2% had dry or very dry skin; while 64.8% considered that their skin was insufficiently hydrated. Mean scores for dryness, burning, itching and stinging/tingling sensation assessed by subjects were 5.3±2.8, 2.4±2.6, 1.5±2.2 and 2.3±2.6, respectively.

Common reasons for dermatologic procedures were skin ageing (49.6%) and acne scars (10.2%)

Subject demographics, skin characteristics, and types of procedures and reasons are provided in Table 1. Incidence and severity of clinical signs are given in Table 2.

After 4 weeks of M89 use, clinical signs (erythema, irritation, desquamation) significantly improved (p<0.0001; Figure 1). Skin hydration had significantly increased in 74.1% of subjects (p<0.0001). Patient symptoms of dryness, burning, pruritus, and stinging/tingling significantly improved as well (p<0.0001; Figure 2).

At study end, 98.4% of subjects were satisfied with the texture of M89. Mean satisfaction score was 8.5 ± 1.7 out of 10 after applying M89 for one week and 9.0 ± 1.5 after 4 weeks. After applying M89 for one week, 93.0% reported that their skin was soothed or very soothed remaining unchanged until week 4. M89 was well-or very well-tolerated by 98.5% of subjects

In 98.0% of subjects, investigator satisfaction was high or very high. In the subgroup treated with "aggressive lasers" (defined as laser resurfacing, laser CO2, Fractional and Erbium lasers; N=99), improvement of clinical signs (Figure 3) and symptoms (Figure 4) was significant (p<0.0001). At study end, all subjects were satisfied with the texture of M89. The mean satisfaction score was 8.7 ± 1.4 out of 10 after applying M89 for one week and 9.1 ± 1.3 after 4 weeks



PREVALENCE OF SUBJECTS WITH CLINICAL SIGNS AT

The difference in prevalence of subjects with improved clinical signs was statistically significant (p<0.0001) after 4 weeks compared to study start





Figure 3

Total n % Erythema 1098 100 Yes 552 50.3 No 546 49.7 Grade 1085 100 Very intense 29 2.7 Intense 83 7.6 Moderate 226 20.8 Low 201 18.5 Absent 546 50.3 Desquamation 1100 100 Yes 594 54.0 No 506 46.0 Grade 1096 100 Very intense 25 2.3 Intense 66 6.0 Moderate 186 17.0 Low 313 28.6 Absent 506 46.2 Irritation 1099 100 Yes 701 63.8 No 398 36.2 Grade 1088 100 Very Intense 22	CLINICAL SIGNS ASSESSED BY THE INVESTIGATORS AT STUDY START			
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Absent 506 46.2 Irritation 1099 100 Yes 701 63.8 No 398 36.2 Grade 1088 100 Very Intense 22 2.0 Intense 82 7.5 Moderate 436 40.1 Low 150 13.8	Moderate	186	17.0	
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Very Intense 22 2.0 Intense 82 7.5 Moderate 436 40.1 Low 150 13.8	No	398	36.2	
Intense 82 7.5 Moderate 436 40.1 Low 150 13.8	Grade	1088	100	
Moderate 436 40.1 Low 150 13.8	Very Intense	22	2.0	
Low 150 13.8	Intense	82	7.5	
	Moderate	436	40.1	
Absent 398 36.6	Low	150	13.8	
	Absent	398	36.6	

PREVALENCE OF SUBJECTS WITH CLINICAL SIGNS AT STUDY START AND AT END OF STUDY AFTER AGGRESSIVE LASER PROCEDURE



The difference in prevalence of subjects with improved clinical signs was statistically significant (p<0.0001) after 4 weeks compared to study start

MEAN CLINICAL SYMPTOM SCORES AT STUDY START AND AT END OF STUDY FOR SUBJECTS WITH SYMPTOMS AT BASELINE HAVING UNDERGONE AGGRESSIVE LASER



The mean score for skin dryness had decreased by 62.1%, for burning sensation by 78.8%, for itching sensation by 70.0% and for stinging/burning tingling by 84.2%. The decrease was statistically significant (p<0.0001).

The mean score for skin dryness had decreased by 70.7%, for burning sensation by 84.9%, for itching sensation by 74.2% and for stinging/ tingling by 91.8%. The decrease was statistically significant (p<0.0001).

CONCLUSION

Daily use of M89 for 4 weeks in subjects having undergone dermatologic procedures, including aggressive lasers resulted in very high user satisfaction along with objective and subjective skin improvement. M89 is an effective and well tolerated adjunct in post-procedure skin care.

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Conflict of interest

Delphine Kerob is an employee of Vichy Laboratoires. Jerry Tan is a member of advisory boards organized by Vichy Laboratoires. The other authors have no conflict of interest to disclose.

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