# A UNIQUE SENSITIVE SKIN CLEANSING TECHNOLOGY

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## BACKGROUND

Sensitive skin is challenging to diagnose since there are different chemical, environmental and psychological triggers and sometimes without visual symptoms. In the first epidemiological study, 51.4% of the women and 38.2% of the men claimed to have sensitive skin [1]. The etiology of sensitive skin is unknown, but some studies suggest the epidermal barrier impairment as a cause of sensitive skin [2]. Shear force and/or friction during cleansing can induce skin damage [3]. Cleansing motion with potential chemical triggers can magnify the symptoms of sensitive skin by inducing barrier damage, yet the skin of afflicted individuals must be maintained in a hygienic condition. In order to improve the skin cleanliness without damage, a novel technology was developed to target sensitive skin individuals.

## OBJECTIVE

- 1. To further the development of a mechanically-moved polymer surface of unique topography to provide treatment and gentle cleansing
- 2. To assess possible differences in treatment/cleansing surface topographies in cleansing sensitive skin.

# **METHODS**

This IRB-approved clinical research study enrolled 16 female subjects, 25-65 years of age, Fitzpatrick skin types I-II, with self-assessed sensitive skin confirmed by the dermatologist investigator in a panel comprised of subjects with rosacea, eczema, atopic dermatitis or cosmetic intolerance syndrome. Subjects used the unique cleansing device constructed to provide a counter-oscillating movement (Figure 1) of a cleansing surface composed of a soft polymer of unique topography and a commercially-available sensitive skin cleanser twice daily for 14 days (Photo 1) Subjects' facial skin was evaluated by self-assessment and clinician grading at baseline and days 7 and 14 both before and after cleansing. A Mann-Whitney two-tailed paired t test was used to analyze the non-parametric data. The device data was compared to baseline longitudinally for the primary treatment/cleansing surface. A separate analysis was conducted for the secondary treatment/cleansing surface for those who experienced irritation with the primary surface.

# **CONFLICT OF INTEREST**

DGK, JN, MR and HEK are employees of Nu Skin Enterprises, Inc. ZDD received funding from Nu Skin Enterprises, Inc.

## RESULTS

The investigator noted immediate improvement following one cleansing in skin smoothness (p=0.009), softness (p=0.017), texture (p=0.028), and cleansing ability (p<0.001). Further sustained improvement occurred in all of these attributes, including pores, with all parameters being highly statistically significant (p<0.001) after 7 days of use with continued excellent performance until the study conclusion at day 14. The sensitive skin subjects noted improvement in skin softness (p=0.011) and smoothness (p=0.008) immediately after one use of the treatment/cleansing device. After 7 days of use, the subjects rated improvement in smoothness (p=0.002), softness (p=0.002), pores (p=0.041), texture (p=0.002), and cleansing ability (p=0.019). This improvement continued into day 14 post cleansing where highly statistically significant (p<0.001) improvement was seen in smoothness, softness, pores, and texture with statistically significant improvement in cleansing ability (p=0.002). All 16 subjects completed the study. No tolerability issues were noted by the investigator dermatologist or the sensitive skin subjects. Additional research is being conducted to evaluate this unique technology.



**Figure 1.** Representation of the counter oscillating motion of the soft-polymer surface.



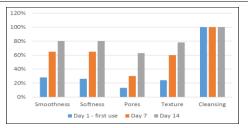
**Photo 1.** Soft-polymer, counter oscillating facial cleansing device.

#### CONCLUSIONS

- A counter-oscillating unique soft polymer cleansing device provided aesthetic improvement and superior cleansing in subjects with sensitive skin.
- Agreement was seen between clinical investigator assessments and subject selfassessments confirming the technological benefit of a novel device-based treatment/cleansing regimen.

#### REFERENCES

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**Figure 2.** Clinical investigator assessment. Percent change over initial visit, untreated, using a 5-point facial attribute grading scale: 0=none, 1=minimal, 2= mild, 3= moderate, 4=severe. Assessments were performed at the research center 10 - 20 minutes after device use in clinic. After Day 7, all improvements were statistically significant (p<0.001).

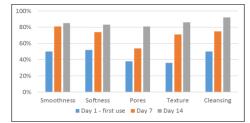


Figure 3. Subject self-assessment. Percent change over initial visit, untreated, using a 5-point facial attribute grading scale: 0=none, 1=minimal, 2= mild, 3= moderate, 4=severe. Assessments were performed at the research center 10 - 20 minutes after device use in clinic. At Day 14, all improvements were statistically significant (p<0.001).