# **Study Report**

A study to evaluate the effectiveness of 
'MEDICUBE AGE-R BOOSTER-H' and one other cosmetic 
product in the skin moisture retention effect 
under dry-weather conditions and 
in the immediate improvement of sleep marks on face

ver.3.0

**December 12, 2022** 



#### **DESCRIPTION FOR SUBMISSION**

An Industrial R&D Center of APR Corporation (hereinafter referred to as "Global Institute of Dermatological Sciences") conducted this study to contribute to the research and development process focusing on the safety and effectiveness of cosmetics sponsored by APR Corporation on the human body. A study to evaluate the effectiveness of 'MEDICUBE AGE-R BOOSTER-H' and one other cosmetic product in the skin moisture retention effect under dry-weather conditions and in the immediate improvement of sleep marks on face was conducted according to following related documents: Guidelines to Application Test to a Human Body and Efficacy Test of Cosmetic Product; Guidelines to Test Methods for the Substantiation Labeling and Advertisement of Cosmetic Product; Guideline for Effectiveness Assessment of Functional Cosmetics by Ministry of Food and Drug Safety; Bioethics and Safety Act by Ministry of Health and Welfare; and Standard Operating Procedure (SOPs) of Global Institute of Dermatological Sciences. The results were reported as follows and this study report is an English translation copy of Korean study report.

#### ver.3.0

#### **December 12, 2022**

### **Global Institute of Dermatological Sciences**



President of Global Institute of Dermatological Sciences:

Dermatologist M.D. Myung Sun Choi

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#### CERTIFICATE OF RELIABILITY ASSURANCE

This study is about the evaluation of the effectiveness of 'MEDICUBE AGE-R BOOSTER-H' and one other cosmetic product in the skin moisture retention effect under dry-weather conditions and in the immediate improvement of sleep marks on face.

This study was appropriately conducted based on the World Medical Association Declaration of Helsinki, and in accordance with following applicable regulatory requirements: "Bioethics and Safety Act\_"; "Cosmetics Act\_" of the Republic of Korea; public announcement from Ministry of Food and Drug Safety; Regulation for Designation of Testing Institutes for Drugs, etc., Cosmetics, and Medical Devices; Korea Good Clinical Practice (KGCP) for Drugs; Guidelines to Application Test to a Human Body and Efficacy Test of Cosmetic Product; Guidelines to Test Methods for the Substantiation Labeling and Advertisement of Cosmetic Product; Guideline on Effectiveness Assessment of Functional Cosmetics; and Standard Operating Procedure (SOPs) of Global Institute of Dermatological Sciences. This report guarantees the reliability of following results.

| Monitoring procedures               | Date                                     | Result   | Note                    |
|-------------------------------------|--|----------|-------------------------|
| Study Protocol                      | October 24, 2022                         | Approved |                         |
| Initial IRB Review                  | October 24, 2022                         | Approved | 70094430-2210-HR-058-03 |
| Study Initiation                    | November 01, 2022                        | A        |                         |
| Study Completion                    | November 02, 2022                        | Approved |                         |
| Data Analysis<br>and Report Work    | November 03, 2022 ~<br>November 10, 2022 | Approved |                         |
| Draft Study Report                  | November 11, 2022                        | Approved |                         |
| English Translation Report          | November 14, 2022                        | Approved | Summary only            |
| IRB Study Closure Report            | November 15, 2022                        | Approved | 70094430-2210-HR-058-07 |
| Final Study Report                  | December 12, 2022                        | Approved |                         |
| Final English Translation<br>Report | December 12, 2022                        | Approved |                         |

Director of Reliability Assurance:

Global Institute of Dermatological Sciences

Hyeyoung Kim



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## **SUMMARY**

| Title of Study                   | A study to evaluate the effectiveness of 'MEDICUBE AGE-R BOOSTER-H' and one other cosmetic product in the skin moisture retention effect under dryweather conditions and in the immediate improvement of sleep marks on face |   |  |
|----------------------------------|--|---|--|
| Study<br>Identification Code     | GLB-221101-I1  |   |  |
| Study<br>Institution             | Global Institute of Dermatological Sciences 4F, 94, Seokchonhosu-ro, Songpa-gu, Seoul, Republic of Korea   |   |  |
| Sponsoring<br>Institution        | APR Corporation 36F, 300, Olympic-ro, Songpa-gu, Seoul, Republic of Korea  |   |  |
| Study<br>Approval Date           | October 24, 2022   |   |  |
| Participation Period of Subjects | November 01, 2022 ~ November 02, 2022  |   |  |
| Reporting Date                   | November 11, 2022  |   |  |
| Subjects                         | 20 Korean adult women aged 20 to 60 who met the inclusion criteria and were not included in the exclusion criteria were enrolled for this study. (Average age: 49.150±6.434_20 completion, 0 drop-out)                       |   |  |
| Study Products                   | Study Product A: At-home skincare device (Industrial product)  'MEDICUBE AGE-R BOOSTER-H'  Study Product B: Basic skin care products, Cream  'MEDICUBE TRIPLE DEEP ERASING CREAM'  |   |  |
| Methods                          | Usage of<br>Study<br>Products  | - Applying study products as a following procedure, once at the first visit after cleansing face and forearm: ① Apply study product B 'MEDICUBE TRIPLE DEEP ERASING CREAM' 0.5 g to the whole face, 0.1 g each to the designated A+B application area (T) and B application area (C1) on left forearm; ② Set study product A 'MEDICUBE AGE-R BOOSTER-H' as Level 5; ③ Apply study product A to the left side of face for 2 minutes, to A+B application area (T) on left forearm for 1 minute. |  |
|                                  | Evaluations  | Effectiveness evaluations     Measurement of skin moisture retention effect under dry weather conditions: Corneometer     Measurement of sleep marks on face: Antera 3D   |  |

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- Safety evaluation
   Subjective questionnaire evaluation
- 1. Results of effectiveness evaluation
- 1) The evaluation results for skin moisture retention effect under dryweather conditions using Corneometer

Significant improvements in skin moisture retention effect under dryweather conditions were observed in comparison to baseline; capacitance, representing hydration of SC (stratum corneum) of epidermis, was increased 65.330% and 56.461% (p<.001) each on A+B application area immediately after once of study products application and after 1 hour exposing under dry-weather conditions, and increased 50.253% and 34.538% (p<.001) each on B application area immediately after once of study product application and after 1 hour exposing under dry-weather conditions; And both application area showed statistically significant (p<.001) differences each to non-application area.

In addition, skin moisture retention effect under dry-weather conditions was further improved when study product A+B were used together; capacitance of A+B application area showed statistically significant (p<.001) difference to B application area immediately after once of study products application and after 1 hour exposing under dry-weather conditions.

Results

2) The evaluation results for immediate improvement of sleep marks on face using Antera 3D

Significant improvements in sleep marks on face were observed on both application area in comparison to baseline (after sleep marks inducement); volume value, representing depressed volume of skin, were decreased 57.964% (p<.001) on A+B application area immediately after once of study products application, and decreased 44.728% (p<.001) on B application area immediately after once of study product application. In addition, sleep marks on face was further improved when study product A+B were used together; volume value of A+B application area showed a statistically significant (p<.01) difference to B application area immediately after once of study products application.

2. Results for safety evaluation

The principal investigator or the sub-investigator entrusted by the principal investigator monitored whether subjects had any of the following skin responses: erythema, edema, scaling, itching, stinging, burning, tightness, and prickling, through asking and assessment. And the

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|            | result showed that abnormal skin responses to application area before and after study products application were not detected during the study period.  |  |  |
|------------|--|--|--|
|            | 3. Results for subjective questionnaire evaluation   |  |  |
|            | Subjective questionnaire was conducted 16 questions: 2 questions about subjects' facial and body skin type, 2 questions about an extent of skin condition improvement after study products application, and 12 questions about satisfaction about study products. And results for each question were taken statistics as average, standard deviation (SD), and positive answer rate (%). |  |  |
| Conclusion | The study products 'MEDICUBE AGE-R BOOSTER-H' and one other cosmetic product, requested by the APR Corporation, were considered to be effective to skin moisture retention effect under dry-weather conditions and to immediately improve sleep marks on face.   |  |  |