

TEST REPORT

2026CS0717

DATE OF RECEPTION

Date Format: dd/MM/yyyy 16/03/2026

DATE TESTS

Starting: 23/03/2026

Ending: 25/03/2026

APPLICANT

KONE LABS SA DE CV
RIO POTOMAC 153 CASTRO DEL RÍO PARQUE
TECNOINDUSTRIAL
MX-36814 Irapuato
Mexico

Att. Amy Rodríguez Rodríguez

IDENTIFICATION AND DESCRIPTION OF SAMPLES

Reference by AITEX	Reference by customer	AITEX sample description
2026CS0717-S01	ÁUREA-L UV Defense Stick	Emulsion/cream in sample collection container

TESTS CARRIED OUT

- SECURITY STUDY FOR THE VERIFICATION OF THE SKIN COMPATIBILITY OF A COSMETIC PRODUCT.



DESCRIPTION OF SAMPLES



Reference by AITEC: 2026CS0717-S01

Reference by customer:

ÁUREA·L UV Defense Stick

AITEC sample description:

Emulsion/cream in sample collection container



RESULTS

SECURITY STUDY FOR THE VERIFICATION OF THE SKIN COMPATIBILITY OF A COSMETIC PRODUCT

1. OBJECTIVE AND PRINCIPLE OF THE STUDY.

Verify the skin compatibility of the cosmetic product referenced as 2026CS0717-S01 after a single application on the skin and under an occlusive patch for 48 hours and under follow-up by the dermatologist.

The reference used has been:

1. Kanto H, Washizaki K, Ito M, Matsunaga K, Akamatsu H, Kawai K, Katoh N, Natsuaki M, Yoshimura I, Kojima H, Okamoto Y, Okuda M, Kuwahara H, Sugiyama M, Kinoshita S, Mori F. Optimal patch application time in the evaluation of skin irritation. J Dermatol. May 2013; 40(5):363-9. doi: 10.1111/1346-8138.12004. Epub February 18, 2013. PMID: 23414058.

2. TYPE OF STUDY

Skin compatibility study under occlusive patch.

3. RESEARCH CENTER AND TECHNICAL TEAM

AITEX

TResearcher: Joaquín Espiñeira Sicre (Dermatologist) Collegiate number: 03/0312448

4. DATE TEST

23/03/2026 - 25/03/2026

5. TESTED PRODUCT REFERENCE

- Reference: 2026CS0717-S01

6. RESULTS

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The experimental conditions have been the following:

- Application area: Back
- Type of patch: Finn Chambers Aqua
- Conditions: Application conditions: Product applied on the aluminum dome.
- Quantity: 50 mm²
- Contact time: 48 h
- Reading: Approximately 30 minutes after removing the patch.

From the results obtained in the experimental conditions adopted, as well as the subjective self-evaluation, the sample referenced as 2026CS0717-S01 presents very good cutaneous acceptance and very good skin tolerance.

VOLUNTEERS

NUMBER

The number of volunteers whose data have been treated at the end of the trial has been 10. Each of them read, understood and signed an Authorization and Commitment Form

INCLUSION CRITERIA

- Number of volunteers: 10
- Age: 18-65 years
- Gender: Both
- Phototype: II-III-IV

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Volunteer information table

N° VOLUNTEER	GENDER	AGE	TYPE OF SKIN	SKIN PHOTOTYPE
1	Male	54	Normal	II
2	Male	24	Normal	III
3	Male	52	Normal	II
4	Female	26	Normal	II
5	Male	28	Normal	II
6	Male	30	Normal	II
7	Female	29	Normal	III
8	Female	54	Normal	III
9	Male	32	Normal	III
10	Male	28	Normal	II

CRITERIA OF NOT INCLUSION

The criteria for non-inclusion were the following:

1. Nickel allergy
2. Allergy or reaction to the same type of products
3. Hyper-reactivity of the skin
- Reactivity to the plaster
5. Severe exposure to the sun at any time during the month prior to the study
6. Exposure to the sun or intense UVA during the test period
7. Sauna baths
8. Intensive or regular practice of one or several sports whose practice can create difficulties
9. Treatment with vitamin A or its derivatives at least 3 months before the start of the study
10. Treatment with topical corticosteroids in the experimental area in the eight days prior to the study

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All the volunteers included in this study are aware and exempt from the previous actions, so they are accepted in said study

A maximum of 10 products have been applied, at the same time, on the same volunteer, the environmental conditions at the time of reading were the following

	Temperature	Humidity
Waiting room	22,5 ° C	46 % HR
Consult dermatology	22 ° C	45 % HR

Table of results of the study

CLASSIFICATION OF THE IRRITATION INDEX

Nº VOLUNTEER	NEGATIVE	DOUBT	WEAK ERITEMA	STRONG	EXTREME
1	X				
2	X				
3	X				
4	X				
5	X				
6	X				
7	X				
8	X				
9	X				
10	X				
SCORE	10	0	0	0	0
VALUE I.R.	0,0	0,0	0,0	0,0	0,0

REACTION INDEX

0,0

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RESULTS ANALYSIS

M&M / ICDRG	ADAPTED SCORE	OBSERVED REACTION
	0	Negative
-	1	Doubtful: clear macules or homogeneous erythema, without infiltration.
+	2	Weak: non-vesicular: erythema, infiltration, papules
++	3	Strong: edema, vesicles: erythema, infiltration, papules, discrete vesicles.
+++	4	Extreme: ulcers/blisters: coalescing vesicles/blisters.

As a result of the test, an Irritation Index is obtained using the following formula:

$$IR = \frac{\sum P(-)*0 + \sum P(1)*1 + \sum P(2)*2 + \sum P(3)*3 + \sum P(4)*4}{\sum P}$$

P= Number of subjects

The result is evaluated as follows:

INTERPRETATION OF RESULTS		
IRRITATION INDEX (IR)	RATING (IR)	SKIN COMPATIBILITY
0 = IR	NON-IRRITATING	VERY GOOD
0 < IR < 0,5	NON-IRRITATING	GOOD
0,5 ≤ IR < 1	SLIGHTLY IRRITATING	INTERMEDIATE
1 ≤ IR < 2	IRRITATING	MODERATE
IR ≥ 2	VERY IRRITATING	BAD

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**CONCLUSION**

Of the 10 volunteers who start the study, they have completed 10.

From the results obtained in the experimental conditions adopted, the
product referenced as 2026CS0717-S01

It has

VERY GOOD SKIN COMPATIBILITY

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I, Dr. Joaquín Espiñeira Sicre Collegiate number: 03/0312448, declare that this study has been developed under my medical responsibility and in accordance with standardized work procedures and in the spirit of the principles of Good Clinical Practices and Regulation (EC) No. 1223/2009 on cosmetic products. Therefore, I declare that the product can carry the claim dermatologically tested.

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