PATCH TEST REPORT

Report Nr. A 23/11/0354

Извештај за тест за закрпи

Извештај бр. A 23/11/0354

COSVOCE AYDINLATICI & PARLATICI YÜZ SERUMU

Tested By | Тестирано од

Root International Research & Development Laboratory

Root Cosmetic Inc.

Mirce Acev br. 2, Skopje 1000, North Macedonia

Мирче Ацев бр. 2, Скопје 1000, Кузеј Македонија

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0354

STUDY OBJECTİVE

Assessment of skin tolerance of a cosmetic product after a single application during 48 hours occlusive patch on 10 volunteers.

CUSTOMER INFORMATION

Applicant:	CAHFT SAĞLIK VE KOZMETİK ÜRÜNLERİ TİCARET LTD. ŞTİ.
Adress:	FATİH MAH. SUSAM SOK. NO:16/A SANCAKTEPE İSTANBUL
Related Person:	-
Contact:	-

SAMPLE INFORMATION

Sample Description:	COSVOCE AYDINLATICI & PARLATICI YÜZ SERUMU
Brand:	-
Appearance:	A homogenous liquid
Package:	Replacement – plastic bottle, with dispensing cap.
Amount:	30 mL
Production Date:	-
Expiry Date:	-
Lot / Serial No:	-
Manufacturer:	CAHFT SAĞLIK VE KOZMETİK ÜRÜNLERİ TİCARET LTD. ŞTİ.
Receipt Date:	10.11.2023
Study Period:	10.11.2023 – 17.11.2023

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0354

TEST INFORMATION

Test Method:	Single application - 48 Hours occlusive patch tests
Time(s) Of Assessment:	1 hour after patch removal, then after 24 & 48 hours (72 hours)
Panel:	10 healthy adult volunteers
Application Area:	On the back
Quantity Of Product:	0.1 g
Methodology Abstract:	The tested product in commercial form in an amount of 0.1g was applied to paper flake (Whatmann 3), which were attached with a porous hypoallergenic (surgical) patch on the shoulders on the straight side or on the back. Treatment sites are assessed before the first application of test material (baseline) and after treatment at 30 minutes after patch removal. Negative controls are used to facilitate evaluation. Skin reactions are scored throughout the test by the same experienced assessor, a dermatologist, who made the baseline assessment and under the same lighting source, following a pre-defined scoring scale. The quantification of the skin irritation is given through a numeric scale (erythema, oedema, dryness/desquamation, vesicles). The average irritant score of the product to be tested is calculated from the average of the quotations obtained for each volunteer, allowing to rank the product from "non irritant to very irritant". Characteristics of volunteers and test results are presented in Table 1.
Related Regulations	1. Regulation (EC) No. 1223/2009 of the European Parliament and of the Council of November 30, 2009 regarding cosmetic products. 2. Cosmetic Europe – The Personal Care Associatio (formerly COLIPA) Guidelines “Product test Guidelines for the Assessment of Human Skin Compatibility 1997”

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0354

In the tested group of 10 people, there were no positive reactions, which proves that the tested product does not have irritating or sensitizing properties.

The test results are presented in Table 1.

Volunteer Tested Person Nr.	Age	Sex	Skin Type	Test Results After 24h	Test Results After 48h
1	28	M	N	(-)	(-)
2	28	W	D	(-)	(-)
3	26	W	N	(-)	(-)
4	32	M	D	(-)	(-)
5	34	M	D	(-)	(-)
6	29	W	D	(-)	(-)
7	31	W	N	(-)	(-)
8	27	M	N	(-)	(-)
9	29	W	N	(-)	(-)
10	28	M	N	(-)	(-)

Sex:	Body Skin Type
W: Woman **M:** Man	**N:** Normal, **D:** Dry, **M:** Mixed, **S:** Sensitive, **L:** Normal with a tendency to oily around the seborrheic of the torso

Skin condition assessment by a dermatologist

0 or (-) - no reaction,
1 or (+/-) - weak erythema,
2 or (+) - erythema,
3 or (++) - erythema, papules,
4 or (+++) - erythema, slight edema,
5 or (++++) - erythema, infiltration, vesicles

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0354

IRRITATION INDEX ($X_{śr}$)

The average irritation index is calculated from the sum of the scores for erythema and edema. Based on the irritation index, the product is classified according to the scale.

AVERAGE IRRITATION INDEX ($X_{śr}$)	PRODUCT CLASSIFICATION
$X_{śr} < 0,5$	non-irritating
$0,5 < X_{śr} < 2,0$	slightly irritating
$2,0 < X_{śr} < 5,0$	moderately irritating
$5,0 \leq X_{śr}$	strongly irritating

Average irritation index for tested product: $X_{śr} = 0$, where $X_{śr}$ = (sum of the scores) / (volunteers number)

CONCLUSION

According to the experimental conditions of the study, the test product, can be considered as

NON-IRRITATING regarding its primary skin tolerance.

Дг. Угур КАЈАСИ

Principal Investigator | Dermatologist

Date: 17.11.2023

Дениз ЕЈСИЕР

Co-Investigator | Research Coordinator

Date: 17.11.2023

PATCH TEST REPORT

Report Nr. A 23/11/0350

Извештај за тест за закрпи

Извештај бр. A 23/11/0350

COSVOCE GÖZ ÇEVRESİ SERUM

Tested By | Тестирано од

Root International Research & Development Laboratory

Root Cosmetic Inc.

Mirce Acev br. 2, Skopje 1000, North Macedonia

Мирче Ацев бр. 2, Скопје 1000, Кузеј Македонија

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0350

STUDY OBJECTİVE

Assessment of skin tolerance of a cosmetic product after a single application during 48 hours occlusive patch on 10 volunteers.

CUSTOMER INFORMATION

Applicant:	CAHFT SAĞLIK VE KOZMETİK ÜRÜNLERİ TİCARET LTD. ŞTİ.
Adress:	FATİH MAH. SUSAM SOK. NO:16/A SANCAKTEPE İSTANBUL
Related Person:	-
Contact:	-

SAMPLE INFORMATION

Sample Description:	COSVOCE GÖZ ÇEVRESİ SERUM
Brand:	-
Appearance:	A homogenous liquid
Package:	Replacement – plastic bottle, with dispensing cap.
Amount:	30 mL
Production Date:	-
Expiry Date:	-
Lot / Serial No:	-
Manufacturer:	CAHFT SAĞLIK VE KOZMETİK ÜRÜNLERİ TİCARET LTD. ŞTİ.
Receipt Date:	10.11.2023
Study Period:	10.11.2023 – 17.11.2023

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0350

TEST INFORMATION

Test Method:	Single application - 48 Hours occlusive patch tests
Time(s) Of Assessment:	1 hour after patch removal, then after 24 & 48 hours (72 hours)
Panel:	10 healthy adult volunteers
Application Area:	On the back
Quantity Of Product:	0.1 g
Methodology Abstract:	The tested product in commercial form in an amount of 0.1g was applied to paper flake (Whatmann 3), which were attached with a porous hypoallergenic (surgical) patch on the shoulders on the straight side or on the back. Treatment sites are assessed before the first application of test material (baseline) and after treatment at 30 minutes after patch removal. Negative controls are used to facilitate evaluation. Skin reactions are scored throughout the test by the same experienced assessor, a dermatologist, who made the baseline assessment and under the same lighting source, following a pre-defined scoring scale. The quantification of the skin irritation is given through a numeric scale (erythema, oedema, dryness/desquamation, vesicles). The average irritant score of the product to be tested is calculated from the average of the quotations obtained for each volunteer, allowing to rank the product from "non irritant to very irritant". Characteristics of volunteers and test results are presented in Table 1.
Related Regulations	1. Regulation (EC) No. 1223/2009 of the European Parliament and of the Council of November 30, 2009 regarding cosmetic products. 2. Cosmetic Europe – The Personal Care Associatio (formerly COLIPA) Guidelines “Product test Guidelines for the Assessment of Human Skin Compatibility 1997”

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0350

In the tested group of 10 people, there were no positive reactions, which proves that the tested product does not have irritating or sensitizing properties.

The test results are presented in Table 1.

Volunteer Tested Person Nr.	Age	Sex	Skin Type	Test Results After 24h	Test Results After 48h
1	28	M	N	(-)	(-)
2	28	W	D	(-)	(-)
3	26	W	N	(-)	(-)
4	32	M	D	(-)	(-)
5	34	M	D	(-)	(-)
6	29	W	D	(-)	(-)
7	31	W	N	(-)	(-)
8	27	M	N	(-)	(-)
9	29	W	N	(-)	(-)
10	28	M	N	(-)	(-)

Sex:	Body Skin Type
W: Woman **M:** Man	**N:** Normal, **D:** Dry, **M:** Mixed, **S:** Sensitive, **L:** Normal with a tendency to oily around the seborrheic of the torso

Skin condition assessment by a dermatologist
0 or (-) - no reaction, 1 or (+/-) - weak erythema, 2 or (+) - erythema, 3 or (++) - erythema, papules, 4 or (+++) - erythema, slight edema, 5 or (++++) - erythema, infiltration, vesicles

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0350

IRRITATION INDEX ($X_{śr}$)

The average irritation index is calculated from the sum of the scores for erythema and edema. Based on the irritation index, the product is classified according to the scale.

AVERAGE IRRITATION INDEX ($X_{śr}$)	PRODUCT CLASSIFICATION
$X_{śr} < 0,5$	non-irritating
$0,5 < X_{śr} < 2,0$	slightly irritating
$2,0 < X_{śr} < 5,0$	moderately irritating
$5,0 \leq X_{śr}$	strongly irritating

Average irritation index for tested product: $X_{śr} = 0$, where $X_{śr}$ = (sum of the scores) / (volunteers number)

CONCLUSION

According to the experimental conditions of the study, the test product, can be considered as

NON-IRRITATING regarding its primary skin tolerance.

Дг. Угур КАЈАСИ

Principal Investigator | Dermatologist

Date: 17.11.2023

Дениз ЕЈСИЕР

Co-Investigator | Research Coordinator

Date: 17.11.2023

PATCH TEST REPORT

Report Nr. A 23/11/0353

Извештај за тест за закрпи

Извештај бр. A 23/11/0353

COSVOCE GÖZENEK SIKILAŞTIRICI SERUM

Tested By | Тестирано од

Root International Research & Development Laboratory

Root Cosmetic Inc.

Mirce Acev br. 2, Skopje 1000, North Macedonia

Мирче Ацев бр. 2, Скопје 1000, Кузеј Македонија

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0353

STUDY OBJECTİVE

Assessment of skin tolerance of a cosmetic product after a single application during 48 hours occlusive patch on 10 volunteers.

CUSTOMER INFORMATION

Applicant:	CAHFT SAĞLIK VE KOZMETİK ÜRÜNLERİ TİCARET LTD. ŞTİ.
Adress:	FATİH MAH. SUSAM SOK. NO:16/A SANCAKTEPE İSTANBUL
Related Person:	-
Contact:	-

SAMPLE INFORMATION

Sample Description:	COSVOCE GÖZENEK SIKILAŞTIRICI SERUM
Brand:	-
Appearance:	A homogenous liquid
Package:	Replacement – plastic bottle, with dispensing cap.
Amount:	30 mL
Production Date:	-
Expiry Date:	-
Lot / Serial No:	-
Manufacturer:	CAHFT SAĞLIK VE KOZMETİK ÜRÜNLERİ TİCARET LTD. ŞTİ.
Receipt Date:	10.11.2023
Study Period:	10.11.2023 – 17.11.2023

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0353

TEST INFORMATION

Test Method:	Single application - 48 Hours occlusive patch tests
Time(s) Of Assessment:	1 hour after patch removal, then after 24 & 48 hours (72 hours)
Panel:	10 healthy adult volunteers
Application Area:	On the back
Quantity Of Product:	0.1 g
Methodology Abstract:	The tested product in commercial form in an amount of 0.1g was applied to paper flake (Whatmann 3), which were attached with a porous hypoallergenic (surgical) patch on the shoulders on the straight side or on the back. Treatment sites are assessed before the first application of test material (baseline) and after treatment at 30 minutes after patch removal. Negative controls are used to facilitate evaluation. Skin reactions are scored throughout the test by the same experienced assessor, a dermatologist, who made the baseline assessment and under the same lighting source, following a pre-defined scoring scale. The quantification of the skin irritation is given through a numeric scale (erythema, oedema, dryness/desquamation, vesicles). The average irritant score of the product to be tested is calculated from the average of the quotations obtained for each volunteer, allowing to rank the product from "non irritant to very irritant". Characteristics of volunteers and test results are presented in Table 1.
Related Regulations	1. Regulation (EC) No. 1223/2009 of the European Parliament and of the Council of November 30, 2009 regarding cosmetic products. 2. Cosmetic Europe – The Personal Care Associatio (formerly COLIPA) Guidelines “Product test Guidelines for the Assessment of Human Skin Compatibility 1997”

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0353

In the tested group of 10 people, there were no positive reactions, which proves that the tested product does not have irritating or sensitizing properties.

The test results are presented in Table 1.

Volunteer Tested Person Nr.	Age	Sex	Skin Type	Test Results After 24h	Test Results After 48h
1	28	M	N	(-)	(-)
2	28	W	D	(-)	(-)
3	26	W	N	(-)	(-)
4	32	M	D	(-)	(-)
5	34	M	D	(-)	(-)
6	29	W	D	(-)	(-)
7	31	W	N	(-)	(-)
8	27	M	N	(-)	(-)
9	29	W	N	(-)	(-)
10	28	M	N	(-)	(-)

Sex:	Body Skin Type
W: Woman **M:** Man	**N:** Normal, **D:** Dry, **M:** Mixed, **S:** Sensitive, **L:** Normal with a tendency to oily around the seborrheic of the torso

Skin condition assessment by a dermatologist
0 or (-) - no reaction, 1 or (+/-) - weak erythema, 2 or (+) - erythema, 3 or (++) - erythema, papules, 4 or (+++) - erythema, slight edema, 5 or (++++) - erythema, infiltration, vesicles

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0353

IRRITATION INDEX ($X_{śr}$)

The average irritation index is calculated from the sum of the scores for erythema and edema. Based on the irritation index, the product is classified according to the scale.

AVERAGE IRRITATION INDEX ($X_{śr}$)	PRODUCT CLASSIFICATION
$X_{śr} < 0,5$	non-irritating
$0,5 < X_{śr} < 2,0$	slightly irritating
$2,0 < X_{śr} < 5,0$	moderately irritating
$5,0 \leq X_{śr}$	strongly irritating

Average irritation index for tested product: $X_{śr} = 0$, where $X_{śr}$ = (sum of the scores) / (volunteers number)

CONCLUSION

According to the experimental conditions of the study, the test product, can be considered as

NON-IRRITATING regarding its primary skin tolerance.

Дг. Угур КАЈАСИ

Principal Investigator | Dermatologist

Date: 17.11.2023

Дениз ЕЈСИЕР

Co-Investigator | Research Coordinator

Date: 17.11.2023

PATCH TEST REPORT

Report Nr. A 23/11/0355

Извештај за тест за закрпи

Извештај бр. A 23/11/0355

COSVOCE KIRMIZI PEELİNG SERUM

Tested By | Тестирано од

Root International Research & Development Laboratory

Root Cosmetic Inc.

Mirce Acev br. 2, Skopje 1000, North Macedonia

Мирче Ацев бр. 2, Скопје 1000, Кузеј Македонија

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0355

STUDY OBJECTİVE

Assessment of skin tolerance of a cosmetic product after a single application during 48 hours occlusive patch on 10 volunteers.

CUSTOMER INFORMATION

Applicant:	CAHFT SAĞLIK VE KOZMETİK ÜRÜNLERİ TİCARET LTD. ŞTİ.
Adress:	FATİH MAH. SUSAM SOK. NO:16/A SANCAKTEPE İSTANBUL
Related Person:	-
Contact:	-

SAMPLE INFORMATION

Sample Description:	COSVOCE KIRMIZI PEELİNG SERUM
Brand:	-
Appearance:	A homogenous liquid
Package:	Replacement – plastic bottle, with dispensing cap.
Amount:	30 mL
Production Date:	-
Expiry Date:	-
Lot / Serial No:	-
Manufacturer:	CAHFT SAĞLIK VE KOZMETİK ÜRÜNLERİ TİCARET LTD. ŞTİ.
Receipt Date:	10.11.2023
Study Period:	10.11.2023 – 17.11.2023

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0355

TEST INFORMATION

Test Method:	Single application - 48 Hours occlusive patch tests
Time(s) Of Assessment:	1 hour after patch removal, then after 24 & 48 hours (72 hours)
Panel:	10 healthy adult volunteers
Application Area:	On the back
Quantity Of Product:	0.1 g
Methodology Abstract:	The tested product in commercial form in an amount of 0.1g was applied to paper flake (Whatmann 3), which were attached with a porous hypoallergenic (surgical) patch on the shoulders on the straight side or on the back. Treatment sites are assessed before the first application of test material (baseline) and after treatment at 30 minutes after patch removal. Negative controls are used to facilitate evaluation. Skin reactions are scored throughout the test by the same experienced assessor, a dermatologist, who made the baseline assessment and under the same lighting source, following a pre-defined scoring scale. The quantification of the skin irritation is given through a numeric scale (erythema, oedema, dryness/desquamation, vesicles). The average irritant score of the product to be tested is calculated from the average of the quotations obtained for each volunteer, allowing to rank the product from "non irritant to very irritant". Characteristics of volunteers and test results are presented in Table 1.
Related Regulations	1. Regulation (EC) No. 1223/2009 of the European Parliament and of the Council of November 30, 2009 regarding cosmetic products. 2. Cosmetic Europe – The Personal Care Associatio (formerly COLIPA) Guidelines “Product test Guidelines for the Assessment of Human Skin Compatibility 1997”

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0355

In the tested group of 10 people, there were no positive reactions, which proves that the tested product does not have irritating or sensitizing properties.

The test results are presented in Table 1.

Volunteer Tested Person Nr.	Age	Sex	Skin Type	Test Results After 24h	Test Results After 48h
1	28	M	N	(-)	(-)
2	28	W	D	(-)	(-)
3	26	W	N	(-)	(-)
4	32	M	D	(-)	(-)
5	34	M	D	(-)	(-)
6	29	W	D	(-)	(-)
7	31	W	N	(-)	(-)
8	27	M	N	(-)	(-)
9	29	W	N	(-)	(-)
10	28	M	N	(-)	(-)

Sex:	Body Skin Type
W: Woman **M:** Man	**N:** Normal, **D:** Dry, **M:** Mixed, **S:** Sensitive, **L:** Normal with a tendency to oily around the seborrheic of the torso

Skin condition assessment by a dermatologist

0 or (-) - no reaction,
1 or (+/-) - weak erythema,
2 or (+) - erythema,
3 or (++) - erythema, papules,
4 or (+++) - erythema, slight edema,
5 or (++++) - erythema, infiltration, vesicles

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0355

IRRITATION INDEX ($X_{śr}$)

The average irritation index is calculated from the sum of the scores for erythema and edema. Based on the irritation index, the product is classified according to the scale.

AVERAGE IRRITATION INDEX ($X_{śr}$)	PRODUCT CLASSIFICATION
$X_{śr} < 0,5$	non-irritating
$0,5 < X_{śr} < 2,0$	slightly irritating
$2,0 < X_{śr} < 5,0$	moderately irritating
$5,0 \leq X_{śr}$	strongly irritating

Average irritation index for tested product: $X_{śr} = 0$, where $X_{śr}$ = (sum of the scores) / (volunteers number)

CONCLUSION

According to the experimental conditions of the study, the test product, can be considered as

NON-IRRITATING regarding its primary skin tolerance.

Дг. Угур КАЈАСИ

Principal Investigator | Dermatologist

Date: 17.11.2023

Дениз ЕЈСИЕР

Co-Investigator | Research Coordinator

Date: 17.11.2023

PATCH TEST REPORT

Report Nr. A 23/11/0351

Извештај за тест за закрпи

Извештај бр. A 23/11/0351

COSVOCE NEMLENDİRİCİ & CANLANDIRICI YÜZ SERUMU

Tested By | Тестирано од

Root International Research & Development Laboratory

Root Cosmetic Inc.

Mirce Acev br. 2, Skopje 1000, North Macedonia

Мирче Ацев бр. 2, Скопје 1000, Кузеј Македонија

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0351

STUDY OBJECTİVE

Assessment of skin tolerance of a cosmetic product after a single application during 48 hours occlusive patch on 10 volunteers.

CUSTOMER INFORMATION

Applicant:	CAHFT SAĞLIK VE KOZMETİK ÜRÜNLERİ TİCARET LTD. ŞTİ.
Adress:	FATİH MAH. SUSAM SOK. NO:16/A SANCAKTEPE İSTANBUL
Related Person:	-
Contact:	-

SAMPLE INFORMATION

Sample Description:	COSVOCE NEMLENDİRİCİ & CANLANDIRICI YÜZ SERUMU
Brand:	-
Appearance:	A homogenous liquid
Package:	Replacement – plastic bottle, with dispensing cap.
Amount:	30 mL
Production Date:	-
Expiry Date:	-
Lot / Serial No:	-
Manufacturer:	CAHFT SAĞLIK VE KOZMETİK ÜRÜNLERİ TİCARET LTD. ŞTİ.
Receipt Date:	10.11.2023
Study Period:	10.11.2023 – 17.11.2023

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0351

TEST INFORMATION

Test Method:	Single application - 48 Hours occlusive patch tests
Time(s) Of Assessment:	1 hour after patch removal, then after 24 & 48 hours (72 hours)
Panel:	10 healthy adult volunteers
Application Area:	On the back
Quantity Of Product:	0.1 g
Methodology Abstract:	The tested product in commercial form in an amount of 0.1g was applied to paper flake (Whatmann 3), which were attached with a porous hypoallergenic (surgical) patch on the shoulders on the straight side or on the back. Treatment sites are assessed before the first application of test material (baseline) and after treatment at 30 minutes after patch removal. Negative controls are used to facilitate evaluation. Skin reactions are scored throughout the test by the same experienced assessor, a dermatologist, who made the baseline assessment and under the same lighting source, following a pre-defined scoring scale. The quantification of the skin irritation is given through a numeric scale (erythema, oedema, dryness/desquamation, vesicles). The average irritant score of the product to be tested is calculated from the average of the quotations obtained for each volunteer, allowing to rank the product from "non irritant to very irritant". Characteristics of volunteers and test results are presented in Table 1.
Related Regulations	1. Regulation (EC) No. 1223/2009 of the European Parliament and of the Council of November 30, 2009 regarding cosmetic products. 2. Cosmetic Europe – The Personal Care Associatio (formerly COLIPA) Guidelines “Product test Guidelines for the Assessment of Human Skin Compatibility 1997”

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0351

In the tested group of 10 people, there were no positive reactions, which proves that the tested product does not have irritating or sensitizing properties.

The test results are presented in Table 1.

Volunteer Tested Person Nr.	Age	Sex	Skin Type	Test Results After 24h	Test Results After 48h
1	28	M	N	(-)	(-)
2	28	W	D	(-)	(-)
3	26	W	N	(-)	(-)
4	32	M	D	(-)	(-)
5	34	M	D	(-)	(-)
6	29	W	D	(-)	(-)
7	31	W	N	(-)	(-)
8	27	M	N	(-)	(-)
9	29	W	N	(-)	(-)
10	28	M	N	(-)	(-)

Sex:	Body Skin Type
W: Woman **M:** Man	**N:** Normal, **D:** Dry, **M:** Mixed, **S:** Sensitive, **L:** Normal with a tendency to oily around the seborrheic of the torso

Skin condition assessment by a dermatologist

0 or (-) - no reaction,
1 or (+/-) - weak erythema,
2 or (+) - erythema,
3 or (++) - erythema, papules,
4 or (+++) - erythema, slight edema,
5 or (++++) - erythema, infiltration, vesicles

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0351

IRRITATION INDEX ($X_{śr}$)

The average irritation index is calculated from the sum of the scores for erythema and edema. Based on the irritation index, the product is classified according to the scale.

AVERAGE IRRITATION INDEX ($X_{śr}$)	PRODUCT CLASSIFICATION
$X_{śr} < 0,5$	non-irritating
$0,5 < X_{śr} < 2,0$	slightly irritating
$2,0 < X_{śr} < 5,0$	moderately irritating
$5,0 \leq X_{śr}$	strongly irritating

Average irritation index for tested product: $X_{śr} = 0$, where $X_{śr}$ = (sum of the scores) / (volunteers number)

CONCLUSION

According to the experimental conditions of the study, the test product, can be considered as

NON-IRRITATING regarding its primary skin tolerance.

Дг. Угур КАЈАСИ

Principal Investigator | Dermatologist

Date: 17.11.2023

Дениз ЕЈСИЕР

Co-Investigator | Research Coordinator

Date: 17.11.2023

PATCH TEST REPORT

Report Nr. A 23/11/0356

Извештај за тест за закрпи

Извештај бр. A 23/11/0356

COSVOCE SEBUM & DENGELEYİCİ GÖZENEK ARINDICIRI YÜZ TONİĞİ

Tested By | Тестирано од

Root International Research & Development Laboratory

Root Cosmetic Inc.

Mirce Acev br. 2, Skopje 1000, North Macedonia

Мирче Ацев бр. 2, Скопје 1000, Кузеј Македонија

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0356

STUDY OBJECTİVE

Assessment of skin tolerance of a cosmetic product after a single application during 48 hours occlusive patch on 10 volunteers.

CUSTOMER INFORMATION

Applicant:	CAHFT SAĞLIK VE KOZMETİK ÜRÜNLERİ TİCARET LTD. ŞTİ.
Adress:	FATİH MAH. SUSAM SOK. NO:16/A SANCAKTEPE İSTANBUL
Related Person:	-
Contact:	-

SAMPLE INFORMATION

Sample Description:	COSVOCE SEBUM & DENGELEYİCİ GÖZENEK ARINDICIRI YÜZ TONİĞİ
Brand:	-
Appearance:	A homogenous liquid
Package:	Replacement – plastic bottle, with dispensing cap.
Amount:	200 mL
Production Date:	-
Expiry Date:	-
Lot / Serial No:	-
Manufacturer:	CAHFT SAĞLIK VE KOZMETİK ÜRÜNLERİ TİCARET LTD. ŞTİ.
Receipt Date:	10.11.2023
Study Period:	10.11.2023 – 17.11.2023

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0356

TEST INFORMATION

Test Method:	Single application - 48 Hours occlusive patch tests
Time(s) Of Assessment:	1 hour after patch removal, then after 24 & 48 hours (72 hours)
Panel:	10 healthy adult volunteers
Application Area:	On the back
Quantity Of Product:	0.1 g
Methodology Abstract:	The tested product in commercial form in an amount of 0.1g was applied to paper flake (Whatmann 3), which were attached with a porous hypoallergenic (surgical) patch on the shoulders on the straight side or on the back. Treatment sites are assessed before the first application of test material (baseline) and after treatment at 30 minutes after patch removal. Negative controls are used to facilitate evaluation. Skin reactions are scored throughout the test by the same experienced assessor, a dermatologist, who made the baseline assessment and under the same lighting source, following a pre-defined scoring scale. The quantification of the skin irritation is given through a numeric scale (erythema, oedema, dryness/desquamation, vesicles). The average irritant score of the product to be tested is calculated from the average of the quotations obtained for each volunteer, allowing to rank the product from "non irritant to very irritant". Characteristics of volunteers and test results are presented in Table 1.
Related Regulations	1. Regulation (EC) No. 1223/2009 of the European Parliament and of the Council of November 30, 2009 regarding cosmetic products. 2. Cosmetic Europe – The Personal Care Associatio (formerly COLIPA) Guidelines “Product test Guidelines for the Assessment of Human Skin Compatibility 1997”

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0356

In the tested group of 10 people, there were no positive reactions, which proves that the tested product does not have irritating or sensitizing properties.

The test results are presented in Table 1.

Volunteer Tested Person Nr.	Age	Sex	Skin Type	Test Results After 24h	Test Results After 48h
1	28	M	N	(-)	(-)
2	28	W	D	(-)	(-)
3	26	W	N	(-)	(-)
4	32	M	D	(-)	(-)
5	34	M	D	(-)	(-)
6	29	W	D	(-)	(-)
7	31	W	N	(-)	(-)
8	27	M	N	(-)	(-)
9	29	W	N	(-)	(-)
10	28	M	N	(-)	(-)

Sex:	Body Skin Type
W: Woman **M:** Man	**N:** Normal, **D:** Dry, **M:** Mixed, **S:** Sensitive, **L:** Normal with a tendency to oily around the seborrheic of the torso

Skin condition assessment by a dermatologist
0 or (-) - no reaction, 1 or (+/-) - weak erythema, 2 or (+) - erythema, 3 or (++) - erythema, papules, 4 or (+++) - erythema, slight edema, 5 or (++++) - erythema, infiltration, vesicles

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0356

IRRITATION INDEX ($X_{śr}$)

The average irritation index is calculated from the sum of the scores for erythema and edema. Based on the irritation index, the product is classified according to the scale.

AVERAGE IRRITATION INDEX ($X_{śr}$)	PRODUCT CLASSIFICATION
$X_{śr} < 0,5$	non-irritating
$0,5 < X_{śr} < 2,0$	slightly irritating
$2,0 < X_{śr} < 5,0$	moderately irritating
$5,0 \leq X_{śr}$	strongly irritating

Average irritation index for tested product: $X_{śr} = 0$, where $X_{śr}$ = (sum of the scores) / (volunteers number)

CONCLUSION

According to the experimental conditions of the study, the test product, can be considered as

NON-IRRITATING regarding its primary skin tolerance.

Дг. Угур КАЈАСИ

Principal Investigator | Dermatologist

Date: 17.11.2023

Дениз ЕЈСИЕР

Co-Investigator | Research Coordinator

Date: 17.11.2023

PATCH TEST REPORT

Report Nr. A 23/11/0352

Извештај за тест за закрпи

Извештај бр. A 23/11/0352

COSVOCE YAŞLANMA KARŞITI VE ONARICI YÜZ SERUMU

Tested By | Тестирано од

Root International Research & Development Laboratory

Root Cosmetic Inc.

Mirce Acev br. 2, Skopje 1000, North Macedonia

Мирче Ацев бр. 2, Скопје 1000, Кузеј Македонија

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0352

STUDY OBJECTİVE

Assessment of skin tolerance of a cosmetic product after a single application during 48 hours occlusive patch on 10 volunteers.

CUSTOMER INFORMATION

Applicant:	CAHFT SAĞLIK VE KOZMETİK ÜRÜNLERİ TİCARET LTD. ŞTİ.
Adress:	FATİH MAH. SUSAM SOK. NO:16/A SANCAKTEPE İSTANBUL
Related Person:	-
Contact:	-

SAMPLE INFORMATION

Sample Description:	COSVOCE YAŞLANMA KARŞITI VE ONARICI YÜZ SERUMU
Brand:	-
Appearance:	A homogenous liquid
Package:	Replacement – plastic bottle, with dispensing cap.
Amount:	30 mL
Production Date:	-
Expiry Date:	-
Lot / Serial No:	-
Manufacturer:	CAHFT SAĞLIK VE KOZMETİK ÜRÜNLERİ TİCARET LTD. ŞTİ.
Receipt Date:	10.11.2023
Study Period:	10.11.2023 – 17.11.2023

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0352

TEST INFORMATION

Test Method:	Single application - 48 Hours occlusive patch tests
Time(s) Of Assessment:	1 hour after patch removal, then after 24 & 48 hours (72 hours)
Panel:	10 healthy adult volunteers
Application Area:	On the back
Quantity Of Product:	0.1 g
Methodology Abstract:	The tested product in commercial form in an amount of 0.1g was applied to paper flake (Whatmann 3), which were attached with a porous hypoallergenic (surgical) patch on the shoulders on the straight side or on the back. Treatment sites are assessed before the first application of test material (baseline) and after treatment at 30 minutes after patch removal. Negative controls are used to facilitate evaluation. Skin reactions are scored throughout the test by the same experienced assessor, a dermatologist, who made the baseline assessment and under the same lighting source, following a pre-defined scoring scale. The quantification of the skin irritation is given through a numeric scale (erythema, oedema, dryness/desquamation, vesicles). The average irritant score of the product to be tested is calculated from the average of the quotations obtained for each volunteer, allowing to rank the product from "non irritant to very irritant". Characteristics of volunteers and test results are presented in Table 1.
Related Regulations	1. Regulation (EC) No. 1223/2009 of the European Parliament and of the Council of November 30, 2009 regarding cosmetic products. 2. Cosmetic Europe – The Personal Care Associatio (formerly COLIPA) Guidelines “Product test Guidelines for the Assessment of Human Skin Compatibility 1997”

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0352

In the tested group of 10 people, there were no positive reactions, which proves that the tested product does not have irritating or sensitizing properties.

The test results are presented in Table 1.

Volunteer Tested Person Nr.	Age	Sex	Skin Type	Test Results After 24h	Test Results After 48h
1	28	M	N	(-)	(-)
2	28	W	D	(-)	(-)
3	26	W	N	(-)	(-)
4	32	M	D	(-)	(-)
5	34	M	D	(-)	(-)
6	29	W	D	(-)	(-)
7	31	W	N	(-)	(-)
8	27	M	N	(-)	(-)
9	29	W	N	(-)	(-)
10	28	M	N	(-)	(-)

Sex:	Body Skin Type
W: Woman **M:** Man	**N:** Normal, **D:** Dry, **M:** Mixed, **S:** Sensitive, **L:** Normal with a tendency to oily around the seborrheic of the torso

Skin condition assessment by a dermatologist

0 or (-) - no reaction,
1 or (+/-) - weak erythema,
2 or (+) - erythema,
3 or (++) - erythema, papules,
4 or (+++) - erythema, slight edema,
5 or (++++) - erythema, infiltration, vesicles

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0352

IRRITATION INDEX ($X_{śr}$)

The average irritation index is calculated from the sum of the scores for erythema and edema. Based on the irritation index, the product is classified according to the scale.

AVERAGE IRRITATION INDEX ($X_{śr}$)	PRODUCT CLASSIFICATION
$X_{śr} < 0,5$	non-irritating
$0,5 < X_{śr} < 2,0$	slightly irritating
$2,0 < X_{śr} < 5,0$	moderately irritating
$5,0 \leq X_{śr}$	strongly irritating

Average irritation index for tested product: $X_{śr} = 0$, where $X_{śr}$ = (sum of the scores) / (volunteers number)

CONCLUSION

According to the experimental conditions of the study, the test product, can be considered as

NON-IRRITATING regarding its primary skin tolerance.

Дг. Угур КАЈАСИ

Principal Investigator | Dermatologist

Date: 17.11.2023

Дениз ЕЈСИЕР

Co-Investigator | Research Coordinator

Date: 17.11.2023

PATCH TEST REPORT

Report Nr. A 23/11/0349

Извештај за тест за закрпи

Извештај бр. A 23/11/0349

NAZİK NEMLENDİRİCİ YÜZ TEMİZLEME YAĞI

Tested By | Тестирано од

Root International Research & Development Laboratory

Root Cosmetic Inc.

Mirce Acev br. 2, Skopje 1000, North Macedonia

Мирче Ацев бр. 2, Скопје 1000, Кузеј Македонија

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0349

STUDY OBJECTİVE

Assessment of skin tolerance of a cosmetic product after a single application during 48 hours occlusive patch on 10 volunteers.

CUSTOMER INFORMATION

Applicant:	CAHFT SAĞLIK VE KOZMETİK ÜRÜNLERİ TİCARET LTD. ŞTİ.
Adress:	FATİH MAH. SUSAM SOK. NO:16/A SANCAKTEPE İSTANBUL
Related Person:	-
Contact:	-

SAMPLE INFORMATION

Sample Description:	Nazik Nemlendirici Yüz Temizleme Yağı
Brand:	-
Appearance:	A homogenous liquid
Package:	Replacement – plastic bottle, with dispensing cap.
Amount:	200 mL
Production Date:	-
Expiry Date:	-
Lot / Serial No:	-
Manufacturer:	CAHFT SAĞLIK VE KOZMETİK ÜRÜNLERİ TİCARET LTD. ŞTİ.
Receipt Date:	10.11.2023
Study Period:	10.11.2023 – 17.11.2023

DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0349

TEST INFORMATION

Test Method:	Single application - 48 Hours occlusive patch tests
Time(s) Of Assessment:	1 hour after patch removal, then after 24 & 48 hours (72 hours)
Panel:	10 healthy adult volunteers
Application Area:	On the back
Quantity Of Product:	0.1 g
Methodology Abstract:	The tested product in commercial form in an amount of 0.1g was applied to paper flake (Whatmann 3), which were attached with a porous hypoallergenic (surgical) patch on the shoulders on the straight side or on the back. Treatment sites are assessed before the first application of test material (baseline) and after treatment at 30 minutes after patch removal. Negative controls are used to facilitate evaluation. Skin reactions are scored throughout the test by the same experienced assessor, a dermatologist, who made the baseline assessment and under the same lighting source, following a pre-defined scoring scale. The quantification of the skin irritation is given through a numeric scale (erythema, oedema, dryness/desquamation, vesicles). The average irritant score of the product to be tested is calculated from the average of the quotations obtained for each volunteer, allowing to rank the product from "non irritant to very irritant". Characteristics of volunteers and test results are presented in Table 1.
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DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0349

In the tested group of 10 people, there were no positive reactions, which proves that the tested product does not have irritating or sensitizing properties.

The test results are presented in Table 1.

Volunteer Tested Person Nr.	Age	Sex	Skin Type	Test Results After 24h	Test Results After 48h
1	28	M	N	(-)	(-)
2	28	W	D	(-)	(-)
3	26	W	N	(-)	(-)
4	32	M	D	(-)	(-)
5	34	M	D	(-)	(-)
6	29	W	D	(-)	(-)
7	31	W	N	(-)	(-)
8	27	M	N	(-)	(-)
9	29	W	N	(-)	(-)
10	28	M	N	(-)	(-)

Sex:	Body Skin Type
W: Woman **M:** Man	**N:** Normal, **D:** Dry, **M:** Mixed, **S:** Sensitive, **L:** Normal with a tendency to oily around the seborrheic of the torso

Skin condition assessment by a dermatologist

0 or (-) - no reaction,
1 or (+/-) - weak erythema,
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4 or (+++) - erythema, slight edema,
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DERMATOLOGICAL TEST REPORT

Report Nr. A 23/11/0349

IRRITATION INDEX ($X_{śr}$)

The average irritation index is calculated from the sum of the scores for erythema and edema. Based on the irritation index, the product is classified according to the scale.

AVERAGE IRRITATION INDEX ($X_{śr}$)	PRODUCT CLASSIFICATION
$X_{śr} < 0,5$	non-irritating
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$5,0 \leq X_{śr}$	strongly irritating

Average irritation index for tested product: $X_{śr} = 0$, where $X_{śr}$ = (sum of the scores) / (volunteers number)

CONCLUSION

According to the experimental conditions of the study, the test product, can be considered as

NON-IRRITATING regarding its primary skin tolerance.

Дг. Угур КАЈАСИ

Principal Investigator | Dermatologist

Date: 17.11.2023

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Co-Investigator | Research Coordinator

Date: 17.11.2023