BİYO TEST LABORATUVARI

Test
TS EN ISO/IEC 17025
AB-1804-T

AB-1804-T
BT-2025-000758
02.25

ANALYSIS REPORT

Purpose of Analysis :	Private Request	**Report Number :**	BT-2025-000758
		Date and Time of Report :	24.02.2025 09:00
Sample requested by:		**Sample Detail:**	
Name	: SERAP DAĞ/ SERAPHY AROMATHERAPY NATURAL SKINCARE	**Name**	: SERAPHY AROMATHERAPY LAVANDER&AMETHYST SALT SCRUB
Address	: SAHİL MAHALLESİ MARMARA CAD. NO : 2 DAİRE 4 ÇİFTLİKKÖY YALOVA	**Qty/Pcs - Temp. (C)**	: 200 mL
		Packing	: Company Packaging
		Date of Prod./Exp.	: 10.02.2025 - 10.02.2027
Authorized Person	: SERAP DAĞ	**Lot Number**	: -
Phone/Fax	:	**Brand**	:
Sender	: SERAP DAĞ/ SERAPHY AROMATHERAPY NATURAL SKINCARE		
Manufacturer	: SERAP DAĞ/ SERAPHY AROMATHERAPY NATURAL	**Date Received**	: 17.02.2025
		Date Started	: 17.02.2025
Offerr No	:	**Date Finished**	: 24.02.2025

RESULT

Name of Analysis	Result	Unit	U	Rec.	LOQ	LVS	D.R.	Reference Ranges	Method/s	Conformity
Detection and Enumeration of Aerobic Mesophilic Bacteria *	<10	cfu/g-ml				1		< 1000	TS EN ISO 21149	Passed
Yeast and Mould *	<10	cfu/g-ml				1		< 1000	TS EN ISO 16212	Passed
Pseudomonas Aeruginosa *	Not Detected.	g-ml				1		Should not be	TS EN ISO 22717	Passed
Staphylococcus Aureus *	Not Detected.	g-ml				1		Should not be	TS EN ISO 22718	Passed
Candida Albicans *	Not Detected.	g-ml				1		Should not be	TS EN ISO 18416	Passed
Escherichia Coli *	Not Detected.	g-ml				1		Should not be	TS EN ISO 21150	Passed

DESCRIPTION

DECISION RULE (D.R.)

:

Limit Value Source (LVS)

1 - Conformity Assessment was carried out according to the 'Guideline on Microbiological Control of Cosmetic Products'.

U. Uncertainty of Measurement **Rec. Recovery** **LOQ. Limit of Quantification**

- When the conformity assessment regarding the test results is given, the regulations, standards, specifications, contracts, etc., if any. The decision rule specified in the documents is used. If there is no decision rule specified in the legislation, the Simple Decision Rule is applied without considering the measurement uncertainty.
-The uncertainties specified in the report are k=2, expanded uncertainty at the 95% confidence interval.
-The results are valid as the sample is received and we are not responsible for the sampling phase. The laboratory cannot be held responsible for the information given by the customer.

REVISION INFORMATION

:

"*" Analysis marked with "*" are within the scope of accreditation.

1. BİYOTEST Laboratory and Consulting Services Ltd., which operates as an Analysis laboratory. Şti. is accredited by TURKAK according to AB-1804-T and TS EN ISO/IEC 17025 standard. The Turkish Accreditation Agency (TURKAK) has signed a Multilateral Agreement with the European Accreditation Association (EA) and a mutual recognition agreement with the International Laboratory Accreditation Association (ILAC) on the recognition of test reports.
2. The results of the Analysis are valid for the above-mentioned sample sent to the laboratory by the company/institution/individual.
3. Descriptive information in the test report that affects the validity of the results has been declared by the customer. Our laboratory is not responsible for any losses/legal obligations that may occur due to the accuracy and use of this information.
4. No part of this test report can be used alone or separately, can not be copied, reproduced or published in whole or in part without the written permission of the laboratory.
5. This report cannot be used for advertising purposes, unsigned and unsealed reports are invalid.

Biyotest Laboratuvarları ve Danışmanlık Hizmetleri Ltd. Şti. **Avcılar V.D. 781549767 - Tic. Sic. No: 327386-5**

Gümüşpala Mah. Kaynata Sok. No:2 Kat:6 Avcılar-İstanbul e-Mail : info@biyotestlab.com Phone : (0212) 591 90 92

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Tuğba ÖZKAN

Head of Sample Submission ad Reporting Department

BİYOTEST LABORATUVARLARI VE DANIŞMANLIK HİZMETLERİ LTD. ŞTİ.
Gümüşpala Mah. Kaynata Sok. No:2 Kat:6
Avcılar İSTANBUL
Avcılar V.D. 781 549 9767

Approved
24.02.2025 09:00:00
Sema YUMAK
Biologist
Manager Of Laboratory

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