

AB-1804-T

BT-2025-000771

02.25

ANALYSIS REPORT

Purpose of Analysis : Private Request	Report Number : BT-2025-000771
	Date and Time of Report : 24.02.2025 09:08
Sample requested by:	Sample Detail:
Name : SERAP DAĞ/ SERAPHY AROMATHERAPY NATURAL SKINCARE	Name : SERAPHY AROMATHERAPY SHEA BUTTER
Address : SAHİL MAHALLESİ MARMARA CAD. NO : 2 DAİRE 4 ÇİFTLİKKÖY YALOVA	Qty/Pcs - Temp. (C) : 200 mL
Authorized Person : SERAP DAĞ	Packing : Company Packaging
Phone/Fax :	Date of Prod./Exp. : 10.02.2025 - 10.02.2027
Sender : SERAP DAĞ/ SERAPHY AROMATHERAPY NATURAL SKINCARE	Lot Number : -
Manufacturer : SERAP DAĞ/ SERAPHY AROMATHERAPY NATURAL SKINCARE	Brand :
Offerr No :	Date Received : 17.02.2025
	Date Started : 17.02.2025
	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.

- BIYOTEST Laboratory and Consulting Services Ltd., which operates as an Analysis laboratory. Şti. is accredited by TÜRKAK according to AB-1804-T and TS EN ISO/IEC 17025 standard. The Turkish Accreditation Agency (TÜRKAK) 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.
- The results of the Analysis are valid for the above-mentioned sample sent to the laboratory by the company/institution/individual.
- 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.
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Page 1 / 2

13.09.2021

P03-T08-F01/

Rev.02/15.12.2022

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Tuğba ÖZKAN

**Head of Sample Submission and
Reporting Department**

BIYOTEST LABORATUVARLARI VE
DANIŞMANLIK HİZMETLERİ LTD. ŞTİ.
Gümüşpala Mah. Kaynata Sok. No:2 Kat:6
Avcılar / İSTANBUL
Avcılar V.D. 781549767

Approved
24.02.2025 09:08:00
Sema YUMAK
Biologist
Manager Of Laboratory

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Page 2 / 2
13.09.2021
P03-T08-F01/
Rev.02/15.12.2022