BİYO TEST LABORATUVARI

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

AB-1804-T
BT-2024-003827
06.24

ANALYSIS REPORT

Purpose of Analysis :	Private Request	**Report Number :**	BT-2024-003827
		Date and Time of Report :	28.06.2024 14:37
Sample requested by:		**Sample Detail:**	
Name	: EVLY PHARMA COSMETICS-The Purest Solutions	**Name**	: Hydration Booster Daily Moisturizing Cream 05/2024
Address	: Şerifali Mah. Söyleşi Sok. No:41/A Ümraniye /İSTANBUL	**Qty/Pcs - Temp. (C)**	: 10 g
		Packing	: Company Packaging
		Date of Prod./Exp.	: 05/2024 - 05/2027
Authorized Person	:	**Lot Number**	: - 0456/1
Phone/Fax	:	**Brand**	:
Sender	: EVLY PHARMA COSMETICS-The Purest Solutions		
Manufacturer	: EVLY PHARMA COSMETICS-The Purest	**Date Received**	: 25.06.2024
		Date Started	: 25.06.2024
Offerr No	:	**Date Finished**	: 28.06.2024

RESULT

Name of Analysis	Result	Unit	U	Rec.	LOQ	LVS	D.R.	Reference Ranges	Method/s	Conformity
Arsenic	<0,01	mg/kg			<0,01	2		< 5	In House Metot	Passed
Lead	<0,01	mg/kg			<0,01	2		< 20	In House Metot	Passed
Cadmium	<0,01	mg/kg			<0,01	2		< 5	In House Metot	Passed
Mercury	<0,01	mg/kg			<0,01	2		< 1	In House Metot	Passed
Antimony	<0,01	mg/kg			<0,01	2		< 10	In House Metot	Passed
Color and Physical Condition Detection *	Suitable.	-				1		It should have its own characteristics	TS EN ISO 5492	Passed

DESCRIPTION

DECISION RULE (D.R.)

:

Limit Value Source (LVS)

1 - Evaluated according to product specification.
2 - Compliance has been evaluated according to the Pharmaceuticals and Medical Devices Agency Guideline on Heavy Metal Impurities in 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.
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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
28.06.2024 14:37
Sema YUMAK
Biologist
Manager Of Laboratory

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