

ANALYSIS REPORT

Purpose of Analysis : Private Request	Report Number : BT-2023-000035
Analysis requested by:	Date and Time of Report : 17.01.2023 11:21
Name : BİYBA NATURA KOZMETİK ÜRÜNLERİ SAN. VE TİC. LTD. ŞTİ.	Sample Detail:
Address : ATIFBEY MAH. 67. SOK. NO:33 İÇ KAPI NO:34 GAZİEMİR/İZMİR	Name : Biyba Skin-Firming + Anti-Cellulite Gel Cream
Authorized Person : SEVİL CAN	Qty/Pcs - Temp. (C) : 12 Pcs
Phone/Fax :	Packing : Company Packaging
Sender : BİYBA NATURA KOZMETİK ÜRÜNLERİ SAN. VE TİC. LTD. ŞTİ.	Date of Prod./Exp. : -
Manufacturer : BİYBA NATURA KOZMETİK ÜRÜNLERİ SAN.	Lot Number : -
Offerr No :	Brand :
	Date Received : 11.01.2023
	Date Started : 11.01.2023
	Date Finished : 17.01.2023

RESULT

Name of Analysis	Result	Unit	U	Rec.	LOQ	LVS	D.R.	Reference Ranges	Method/s	Conformity	
Paraben Determination											
Methyl Paraben	Not Detected.	mg/kg						1	Should Not Be	96/45/EC	Passed
Ethyl Paraben	Not Detected.	mg/kg						1	Should Not Be	96/45/EC	Passed
Butyl Paraben	Not Detected.	mg/kg						1	Should Not Be	96/45/EC	Passed
izobütıl paraben	Not Detected.	mg/kg						1	Should Not Be	96/45/EC	Passed
propyl paraben	Not Detected.	mg/kg						1	Should Not Be	96/45/EC	Passed
Isopropyl Paraben	Not Detected.	mg/kg						1	Should Not Be	96/45/EC	Passed
Benzyl Paraben	Not Detected.	mg/kg						1	Should Not Be	96/45/EC	Passed
Hexamidine Paraben	Not Detected.	mg/kg						1	Should Not Be	96/45/EC	Passed
Hexamidine Diparaben	Not Detected.	mg/kg						1	Should Not Be	96/45/EC	Passed
Gluten Determination	Not Detected.	mg/kg						1	Should Not Be	ELISA	Passed
Determination of 1.4 Dioxane	< 0,3	mg/kg			< 0,3					In House Metot	Not Evaluate
Arsenic	< 0,0059	mg/kg			<0,0059	3		< 5		In House Metot	Passed
Lead	< 0,0059	mg/kg			<0,0059	3		< 20		In House Metot	Passed
Cadmium	< 0,0058	mg/kg			<0,0058	3		< 5		In House Metot	Passed
Mercury	< 0,0059	mg/kg			<0,0059	3		< 1		In House Metot	Passed
Antimony	< 0,0058	mg/kg			<0,0058	3		< 10		In House Metot	Passed

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

- BIYOTEST Laboratory and Consulting Services Ltd., which operates as an experiment 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.
- The results of the experiments 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.
- 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.
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Detection and Enumeration of Aerobic Mesophilic Bacteria *	< 10	cfu/g				2		< 1000	TS EN ISO 21149	Passed
Yeast and Mould *	< 10	cfu/g				2		< 1000	TS EN ISO 16212	Passed
Pseudomonas Aeruginosa *	Not Detected.	g				2		Should not be	TS EN ISO 22717	Passed
Staphylococcus Aureus *	Not Detected.	g				2		Should not be	TS EN ISO 22718	Passed
Candida Albicans *	Not Detected.	g				2		Should not be	TS EN ISO 18416	Passed
Escherichia Coli *	Not Detected.	g				2		Should not be	TS EN ISO 21150	Passed
Animal DNA Search (Vegan Test)	Not Detected.					1		Should Not Be	Kit Method,ISO 21571:2005	Passed
Determination of Silicon	Not Detected.	w/w %				1		Should not be	In House Metot	Passed
Paraffin Determination	Not Detected.	%				4		Should not be	In House Metot	Passed

DESCRIPTION

DECISION RULE (D.R.)

:

Limit Value Source (LVS)

- 1 - Evaluation has been made according to the specifications regarding the "does not contain" claims on the customer label.
- 2 - Conformity Assessment was carried out according to the 'Guideline on Microbiological Control of Cosmetic Products'.
- 3 - Compliance has been evaluated according to the Pharmaceuticals and Medical Devices Agency Guideline on Heavy Metal Impurities in Cosmetic Products.
- 4 - Evaluation has been made according to the specifications regarding the "does not contain" claims on the customer label.

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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

Tuğba ÖZKAN

Head of Sample Submission and
Reporting Department

Melda BAĞÇE
Biologist

Head of Microbiology
Department

Pınar Güler GÖKÇE
Chemist

Head of Chemistry
Laboratory Department

BIYOTEST LABORATUVARLARI VE
DANIŞMANLIK HİZMETLERİ LTD. ŞTİ.
Gümüşpala Mah. Kaynata Sok. No:2 Kat:5
Avclar V.D. 81549767
Avclar V.D. 81549767

Approved
17.01.2023 11:21
Sema YUMAK
Biologist
Manager Of Laboratory

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