BİYO TEST LABORATUVARI

TÜRKAK

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

AB-1804-T

BT-2023-000032

01.23

ANALYSIS REPORT

Purpose of Analysis :	Private Request	**Report Number :**	BT-2023-000032
		Date and Time of Report :	17.01.2023 11:14
Analysis requested by:		**Sample Detail:**	
Name	: BİYBA NATURA KOZMETİK ÜRÜNLERİ SAN. VE TİC. LTD. ŞTİ.	**Name**	: Biyba Rosy Pearl Body Spray
Address	: ATIFBEY MAH. 67. SOK. NO:33 İÇ KAPI NO:34 GAZİEMİR/İZMİR	**Qty/Pcs - Temp. (C)**	: 12 Pcs
		Packing	: Company Packaging
		Date of Prod./Exp.	: -
Authorized Person	: SEVİL CAN	**Lot Number**	: -
Phone/Fax	:	**Brand**	:
Sender	: BİYBA NATURA KOZMETİK ÜRÜNLERİ SAN. VE TİC. LTD. ŞTİ.		
Manufacturer	: BİYBA NATURA KOZMETİK ÜRÜNLERİ SAN.	**Date Received**	: 11.01.2023
		Date Started	: 11.01.2023
Offerr No	:	**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
izobütil 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.

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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

ANALYSIS REPORT

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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				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 ad Reporting Department

Melda BAĞÇE
Biologist
Head of Microbiology Department

Pınar Güler GÖKÇE
Chemist
Head of Chemistry Laboratory Department

BİYOTEST LABORATUVARLARI VE DANIŞMANLIK HİZMETLERİ LTD. ŞTİ.
Gümüşpala Mah. Kaynata Sok. No:2 Kat:5
Avcılar / İSTANBUL
Avcılar V.D. 781 154 9767

Approved
17.01.2023 11:14
Sema YUMAK
Biologist
Manager Of Laboratory

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