NANOLAB LABORATORY SERVICES

REPORT of EXAMINATION and ANALYSIS

Test
TS EN ISO/IEC 17025
AB-0566-T

AB-0566-T
U23-2253/0
09-23

Report No / Rev. No : **U23-2253/0** **Report Date** : 08.09.2023
The Purpose of Analysis : SpecialRequest
Sample Sent by : ÇEKA AMBALAJ KAĞ.VE MED.SAN.DIŞ TİC.LTD.ŞTİ.
Address : TURGUT REİS MH.DEMOKRASİ CD.NO:171/2, SULTANBEYLİ/İSTANBUL
Sample Name : PIPETTE
Sample Quantity : 50 ADET/PCS **Sample Package** : -
Temperature (°C) : 25 °C
Sample Acc. Date & Time : 31.08.2023 19:11
Analysis Start - Finish Date : 01.09.2023 - 08.09.2023

Analysis	Results	Method / Device	R (%)	E.U.(±)	LOQ	Limit	E
1-Chloride (%)	0,08	TS EN 645-1996, Mohr Yöntemi Titrimetric Method	94,052	0,03		≤ 0,2	P
2-Formaldehyde (mg/kg)	Not Detected	BS-EN 13130-23, TS EN 645 Spektrofotometrik Metot	94,052		1	≤ 15	P
3-Pentachlorophenol (mg/kg)	Not Detected	ISO 15318 LC-MS/MS	94,052		0,01	≤ 0,15	P
4-*Arsenic (mg/kg)	Not Detected	In-house Method-"K.SOP.212/Rev.01" (Modified from NMKL 186) ICP-MS				≤ 2	P
5-*Cadmium (mg/dm²)	Not Detected	In-house Method-"K.SOP.212/Rev.01" (Modified from NMKL 186) ICP-MS				≤ 0,002	P
6-*Lead (mg/dm²)	Not Detected	In-house Method-"K.SOP.212/Rev.01" (Modified from NMKL 186) ICP-MS				≤ 0,003	P

Gül GÜVENÇ
ERMM Lab.
Manager
e-signed

Bülent TATLISÖZ
Manager of Sample
Accept. and Report
e-signed

Confirmable
08.09.2023
Yunus Emre YILMAZ
Lab. Manager
e-signed

NANOLAB LABORATUVAR HİZMETLERİ NANOLAB MERKEZ

This document has been electronically signed in accordance with the Electronic Signature Law No. 5070.
https://eimza.nano-lab.com.tr/home/Index?privateCode=CDD3DB71&date=08.09.2023

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Analysis	Results	Method / Device	R (%)	E.U.(±)	LOQ	Limit	E
7-Mercury (Hg) (mg/dm²)	Not Detected	In-house Method- "K.SOP.212/Rev.01" (Modified from NMKL 186) ICP-MS	96,00			≤ 0,002	P

Nanolab Laboratuvar Hizmetleri Kimya Gıda Danışmanlık Çevre Eğitim San. ve Tic. Ltd. Şti. accredited by TÜRKAK under registration number AB-0566-T for TS EN ISO / IEC 17025 as test laboratory

Turkish Accreditation Agency (TURKAK) is a signatory to the European co-operation for Accreditation (EA) Multilateral Agreement (MLA) and to the International Laboratory Accreditation Cooperation (ILAC) Mutual Recognition Arrangement (MRA) for the recognition of test reports

1. No part of the analysis report can not be used alone or separately.
2. This report can not be used in judicial-administrative proceedings and for advertising purposes.
3. Analysis results are valid for the above mentioned sample.
4. This report may not be partially copied or reproduced without the written permission of the laboratory.
5. Unsigned and unsealed reports are not valid.
6. The above mentioned values were determined as the result of the examination and analysis.
7. Decision Rule: The conformity statement has been made in favor of the producer using quantitative physical and chemical analyses without considering measurement uncertainty in microbiological, sensory, and qualitative analyses (Simple Acceptance Rule).
8. NI: Within the scope of the relevant legislation, no evaluation can be made for analyzes that do not have a limit value.
9. The analysis signed with "*" are in the scope of accreditation.
10. The laboratory is not responsible for the information declared by the customer.
11. Abbreviations; E : Evaluation, P : Pass, F : Fail, N.I. : Not Interpreted, R : Recovery, E.U. : Expanded Uncertainty, LOQ : Limit of Quantification

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Manager
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Manager of Sample
Accept. and Report
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