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CERTIFICATE OF ANALYSIS No. 20241016-M4651

Product Information		Sample Information	
Product Name:	CBD Suppository 75mg	Sample Received ¹ :	2024-10-16
Product Type:	Suppositories	Sample Condition:	Suitable
Product ID:	SD30722	Analysis Completed:	2024-10-16
Batch No.:	B2979	Certificate Dated:	2024-10-16
Manufacture Date:	2024-10-16	Retest Interval:	24 months

RESULTS

Parameter	Method	Requirement/ Limits	Results	Compliant / Non-Compliant ²
Identification	UV/Vis	N/A	Retention time complies with Certified Reference Material	N/A
Cannabidiol (CBD) %		N/A	80.398 mg	N/A
Additional cannabinoids:				N/A
CBG, %		N/A	2.009 mg	N/A
CBC, %		N/A	2.984 mg	N/A
CBN %		N/A	0.911 mg	N/A
CBDV, %		N/A	1.191 mg	N/A
Total	HPLC			
Tetrahydrocannabinol, %		< 3.64 mg (<0.2%)	3.4 mg (0.183%)	N/A
CBDA, %		N/A	0.372 mg	N/A
CBGA, %		N/A	ND	N/A
THCV, %		N/A	ND	N/A
CBL, %		N/A	0.372 mg	N/A
CBDVA, %		N/A	ND	N/A

THE END OF THE CERTIFICATE

Authorized by: Quality Control Analyst 2024-10-16
Approved by: Senior Quality Control Analyst 2024-10-16

¹ Samples were delivered by the Client.
² UAB Biosyyd applies the acceptance decision rule in accordance with ILAC-G8:09/2019: if measurement plus minus expanded measurement uncertainty meets specification requirements, the parameter is stated as compliant.
³ Given expanded measurement uncertainty was estimated for the coverage factor k = 2 at 95 % confidence level. Sampling uncertainty has not been taken into consideration.
⁴ Test method is accredited.
⁵ Test performed on Cannabis Sativa raw material in Third party laboratory.

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PRODUCT CONTAMINATION REPORT

Product Information			Sample Information	
Product Name:	CBD Suppository 75mg		Sample Received ¹ :	2024-10-16
Product Type:	Suppositories		Sample Condition:	Suitable
Product ID:	SD30722		Analysis Completed:	2024-10-16
Batch No.:	B2979		Certificate Dated:	2024-10-16
Manufacture Date:	2024-10-16		Retest Interval:	24 months
RESULTS				
Parameter	Method	Requirement/ Limits	Results	Compliant / Non-Compliant ²
Residual Solvents:				
Ethanol	Gas Chromatography	≤ 50 ppm	ND	Compliant
n-Pentane	Gas Chromatography	≤ 50 ppm	ND	Compliant
2-propanol	Gas Chromatography	≤ 50 ppm	ND	Compliant
n-Heptane	Gas Chromatography	≤ 50 ppm	ND	Compliant
Microbiological quality ⁵ :				
Enumeration of mesophilic aerobic bacteria	PN-EN ISO 4833-1:2013-12	≤ 10 ⁵ CFU/g	4.0 · 10 ¹ CFU/g	Compliant
Enumeration of yeasts	PN-ISO 21527-2:2009	≤ 10 ³ CFU/g	< 1.0 · 10 ¹ CFU/g	Compliant
Enumeration of moulds	PN-ISO 21527-2:2009	≤ 10 ³ CFU/g	< 1.0 · 10 ¹ CFU/g	Compliant
Specified microorganisms:				
Enumeration of <i>Bacillus Cereus</i>	PN-EN ISO 7932:2005	≤ 10 ³ CFU/g	< 1.0 · 10 ¹ CFU/g	Compliant
Enumeration of coagulase-positive staphylococci (<i>Staphylococcus aureus</i> and other species)	PN-EN ISO 6888-1:2001+A1:2004	≤ 10 ³ CFU/g	< 1.0 · 10 ¹ CFU/g	Compliant
Detection of <i>Salmonella</i> spp.	PN-EN ISO 6579-1:2017-04	Not Detected in 25 g	Not Detected in 25 g	Compliant
Pesticides ⁵ :				
Organochlorine pesticides	LMBG-00.00-34:1999 (DFG S19) except section E9	Limits of each Pesticides according to Ph. Eur 2.8.13	< LOQ mg/kg	Compliant
Organophosphorus pesticides			< LOQ mg/kg	Compliant
Pyrethroids			< LOQ mg/kg	Compliant
Propachlor		≤ 0.1 mg/kg	< LOQ mg/kg	Compliant
Aflatoxin B1 ⁵	AFL/01/2012/1	≤ 5 µg/kg	< 1.0 µg/kg	Compliant
Aflatoxin (sum um or R1+R2+G1+G2) ⁵	AFL/01/2012/1	≤ 10 µg/kg	< LOQ µg/kg	Compliant
Ochratoxin A ⁵	PB-456 ed. I of 15.10.2021	≤ 10 µg/kg	< 0.25 µg/kg	Compliant
Heavy metals ⁵ :				
Arsenic	PN-EN 15763:2010	≤ 1.0 mg/kg	0.13 ± 0.02 mg/kg	Compliant
Cadmium	PN-EN 15763:2010	≤ 0.4 mg/kg	0.018 ± 0.004 mg/kg	Compliant
Lead	PN-EN 15763:2010	≤ 1.0 mg/kg	0.39 ± 0.10 mg/kg	Compliant
Mercury	PN-EN 15763:2010	≤ 0.2 mg/kg	0.012 ± 0.002 mg/kg	Compliant

⁵ Tested in third part laboratory

We declare that this product is free from microbiological and chemical contaminants including residual solvents, microbiological contamination, pesticides, aflatoxins and heavy metals. We ensure this by performing testing on all raw materials used and periodical testing on final products for self control. This report is applicable only to the batch that is written on report. This report cannot be reproduced partially without a prior written consent of UAB Biosyyd, and is restricted exclusively to the results and statements presented in the original copy of the Report.