

Product	Cannabistada Extrakt THC 50/ CBD 25 CA
Country	Germany
License No.	DE_RP_01_MIA_2022_0054
Potency	Δ^9 -Tetrahydrocannabinol: 50 mg/ml Cannabidiol: 25 mg/ml
Dosage form	Oral liquid
Pack size	30 ml amber glass bottle
Batch number:	G23ACK-O4F1

Manufacturer:	MediPharm Labs Inc. 151 John Street, Barrie, Ontario, L4N 2L1 Canada	Manufacturing date: 04/2023
Manufacturing license No.:	LIC-DZIH2YVSJ-2021	
Manufacturer of Cannabis Flos (starting material):	Northern Green Canada Inc., 275 Orenda Rd., Brampton, L6T 3T7 – Ontario, Canada; Emblem Cannabis Corporation, 20 Woodslee Avenue, Paris, N3L 3N6 – Ontario, Canada	
Strain:	Gorilla Glue 4 Cannatonic	

Test	Method	Specification	Result	Complies
Appearance	Organoleptic	Greenish or yellow to brown liquid	Complies	YES
Identification	DAB Monograph (TLC)	Complies with the description of the DAB monograph	Complies	YES
Relative density	EP 2.2.5	For information	0.9541	YES
Water content	EP 2.5.12	≤ 0.5 %	0.12%	YES
Assay	DAB Monograph (HPLC)	CBD: 22.5 mg/ml – 27.5 mg/ml CBN: ≤ 2.5% THC: 45 mg/ml – 55 mg/ml	27.17 mg/ml 0.15 % 53.58 mg/ml	YES
Ethanol Isopropyl Alcohol	EP 2.4.24	≤ 5000 ppm ≤ 5000 ppm	n.d. n.d.*	YES
Pesticides	EP 2.8.13	Complies with the requirements EP 2.8.13	Complies *	YES
Aflatoxins	EP 2.8.18	B1: ≤ 2.0 µg/kg Sum B1, B2, G1, G2: ≤ 4 µg/kg	< 2 µg/kg * < 4 µg/kg *	YES
Heavy metals analysis	EP 2.4.27	Arsenic: ≤ 2.0 ppm Lead: ≤ 5.0 ppm Mercury: ≤ 0.1 ppm Cadmium: ≤ 0.5 ppm	< 0.1 ppm * < 0.1 ppm * < 0.1 ppm * < 0.05 ppm *	YES
Microbiological purity				YS
<i>TAMC</i>	EP 2.6.12	≤ 10 ⁴ cfu/g or cfu/ml	< 10 cfu/g	
<i>TYMC</i>		≤ 10 ² cfu/g or cfu/ml	< 10 cfu/g	
<i>Bile tolerant gram neg. bacteria</i>	EP 2.6.13	≤ 10 ² cfu/g or cfu/ml	< 10 cfu/g	
<i>Escherichia coli</i>		absent / 1 g or 1 ml	absent/ 1g	
<i>Salmonella species</i>		absent / 10 g or 10 ml	absent/ 10g	
<i>Staphylococcus aureus</i>		absent / 1 g or 1 ml	absent/ 1g	

n.d. - below detection limit, Ethanol 0,05%

Examined: WESSLING GmbH, Johann-Krane-Weg 42, 48149 Münster

*CoA MediPharm Labs Canada Inc., Barrie, Canada

**Expiry date**

04/2025

I hereby declare that this batch of the medicinal product has been manufactured and tested in compliance with the applicable GMP regulations and in accordance with the master documents approved by the client and has been tested in compliance with recognized pharmaceutical regulations pursuant to § 6 No. 3 ApBetrO and is hereby released for marketing pursuant § 16 AMWHV.

All starting materials and intermediate products used have been tested and were approved.

Dr. Klaus-Uwe Pechar

22.06.2023

Date / Signature Qualified Person