



Certificate of Analysis CANNABIS Flos

Batch: 13501

Product	Mother 26 GF CAN
Country	Germany
Manufacturing license No.:	DE RP 01 MIA 2022 0054
Potency	Δ^9 -Tetrahydrocannabinol approx. 26% Cannabidiol $\leq$ 1%
Dosage form	Cannabis Flowers, dried; cultivar: Garlic Funk
Pack size	10g metallized PET pouches 100g metallized PET pouches
Batch bulk	1888GF64901
Batch	13501

Manufacturing	ABCann Medicinals Inc., 126 Vanluven Road, Napane, Ontario K7R 3L2, Canada	
Labelling	PS Pharma Service GmbH Lise-Meitner-Str. 10 40670 Meerbusch	Manufacturing Date: 04.04.2025
Manufacturing license No.:	DE NW 03 MIA 2024 0050	

Test	Method	Specification	Result	Complies
Identification A	EP 3028	Complies with the description of the monograph	Complies	YES
Identification B	EP 3028	Complies with the description of the monograph	Complies	YES
Identification C	EP 3028 (TLC)	Complies with the description of the monograph	Complies	YES
Assay	EP 3028 (HPLC)	CBD total: $\leq$ 1.0% CBN: $\leq$ 1.0% THC total: 23.4% – 28.6%	< 0.1 % 0.1% 24.8 %	YES
Foreign matter	EP 2.8.2	$\leq$ 2.0%	< 2 %	YES
Loss on drying	EP 3028	$\leq$ 12.0%	7.4 %	YES
Pesticides*	EP 2.8.13	Complies with the requirements EP 2.8.13	Complies	YES
Aflatoxins*	EP 2.8.18	B1: $\leq$ 2 μ g/kg Sum B1, B2, G1, G2: $\leq$ 4 μ g/kg	n.d. n.d.	YES
Ochratoxin A*	EP 2.8.22	$\leq$ 20 μ g/kg	n.d.	YES
Heavy metals analysis*	EP 2.4.27	Arsenic: $\leq$ 0.2 ppm Cadmium: $\leq$ 0.3 ppm Lead: $\leq$ 0.5 ppm Mercury: $\leq$ 0.1 ppm	< 0.1 ppm < 0.1 ppm < 0.1 ppm < 0.05 ppm	YES
Microbiological purity*		EP 5.1.8C		YES
TAMC	EP 2.6.12	$\leq$ 500 000 cfu/g	< 100 cfu/g	
TYMC		$\leq$ 50 000 cfu/g	< 10 cfu/g	
Bile tolerant gram neg. bacteria	EP 2.6.31	$\leq$ 10 000 cfu/g	< 10 cfu/g	
Escherichia coli		absent / 1 g	absent / g	
Salmonella species		absent / 25 g	absent / 25g	

n.d. = below limit of quantitation

Examined by: QSI GmbH, Flughafendamm 9a, 28199 Bremen

*Eurofins BioPharma Product Testing Toronto, Canada

Expiry date	12/2025
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BIC: MALADE51KOB

GF: Nicole Broockmann



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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.

08.04.2025

Dr. Klaus-Uwe Pechar

Date / Signature Qualified Person

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