



Certificate of Analysis CANNABIS Flos

Batch: 16502

Product	Mother 28 B8 CAN
Country	Germany
Manufacturing license No.:	DE RP 01 MIA 2022 0054
Potency	Δ^9 -Tetrahydrocannabinol approx. 28% Cannabidiol $\leq$ 1%
Dosage form	Cannabis Flowers, dried; cultivar: Beta 808
Pack size	10g metallized PET pouches 100g metallized PET pouches
Batch bulk	1909BA66601
Batch	16502

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: 05.2025
Manufacturing license No.:	DE NW 03 MIA 2025 0019	

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: 25.2% – 30.8%	< 0.1 % < 0.1% 26.8 %	YES
Foreign matter	EP 2.8.2	$\leq$ 2.0%	< 2 %	YES
Loss on drying	EP 3028	$\leq$ 12.0%	9.3 %	YES
Pesticides*	EP 2.8.13	Complies with the requirements EP 2.8.13	n.n.	YES
Aflatoxins*	EP 2.8.18	B1: $\leq$ 2 μ g/kg Sum B1, B2, G1, G2: $\leq$ 4 μ g/kg	n.n. n.n.	YES
Ochratoxin A*	EP 2.8.22	$\leq$ 20 μ g/kg	n.n.	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	n.n. n.n. n.n. n.n.	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.n. = below limit of quantitation

Examined by:

QSI GmbH, Flughafendamm 9a, 28199 Bremen

* Eurofins BioPharma Product Testing Toronto Inc., Canada

Expiry date	01/2026
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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.

 14.05.2025 
Date / Signature Qualified Person

ADREXpharma GmbH, Jakob- Hasslacher- Str. 4 56070 Koblenz
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