

Certificate of Analysis CANNABIS Flos Batch: K.18.DRP00270-1

Product	Bava Medical 21/1 PLA-DO	
Country	Germany	
Manufacturing license No.:	DE_RP_01_MIA_2022_0054	
Potency	Δ9-Tetrahydrocannabinol approx. 21%; Cannabidiol ≤ 1%	
Dosage form	Cannabis flowers, dried; cultivar: Platinum Dosi-Dos	
Pack size	400g pouches	
	5g cans	
Batch	K.18.DRP00270-1	

Manufacturing date:	27.06.2025

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: ≤ 1.0% CBN: ≤ 1.0% THC total: 18.9% – 23.1%	< 0.1% 0.5% 22.8%	YES
Foreign matter	EP 3028	≤ 2.0% No seeds No leaves > 1.0 cm	< 2 % Complies Complies	YES
Loss on drying	EP 3028	≤ 12.0%	6.0%	YES
Total Ash	EP 2.4.16	≤ 20%	9.8%	YES
Pesticides	EP 2.8.13	Complies with the requirements EP 2.8.13	n.n.	YES
Aflatoxins	EP 2.8.18	B1: ≤ 2 μg/kg Sum B1, B2, G1, G2: ≤ 4 μg/kg	n.n. n.n.	YES
Ochratoxin A	EP 2.8.22	≤ 20 µg/kg	n.n.	YES
Heavy metals analysis	EP 2.4.27	Arsenic: ≤ 0.2 ppm Cadmium: ≤ 0.3 ppm Lead: ≤ 0.5 ppm Mercury: ≤ 0.1 ppm	n.n. n.n. n.n. n.n.	YES
Microbiological purity*		EP 5.1.8C		YES
TAMC TYMC Bile tolerant gram neg.	EP 2.6.12	≤ 500 000 cfu/g ≤ 50 000 cfu/g	< 1000 cfu/g < 10 cfu/g	
bacteria Esherichia coli Salmonella species	EP 2.6.31	≤ 10 000 cfu/g absent / 1 g absent / 25 g	< 100 cfu/g absent /g absent /25g	

n.n. = below limit of quantitation

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

Expiry date	30.11.2025
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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.

01.07.2025

Dr. Klaus-Uwe Pechar

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Date / Signature Qualified Person

ADREXpharma GmbH, Jakob- Hasslacher- Str. 4 56070 Koblenz HRB 26108, Koblenz

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