

Product	CannabiStada 22/1 WAP
Country	Germany
Manufacturing license No.:	DE_RP_01_MIA_2022_0054
Potency	Δ^9 -Tetrahydrocannabinol approx. 22%
	Cannabidiol $\leq 1\%$
Dosage form	Cannabis Flowers, dried; cultivar: Wappa
Pack size	10 g PET/PE cans
	25 g metallized PET pouches
Batch	34401

Grower/drier	ABcann Medicinals Inc., 126 Vanluven Road, Napane, Ontario K7R 3L2, Canada	
Packager	PS Pharma Service GmbH Lise-Meitner-Str. 10 40670 Meerbusch	Manufacturing Date: 02.09.2024
Manufacturing license No.:	DE_NW_03_MIA_2024_0023	

Test	Method	Specification	Result	Complies
Odour	Organoleptic	Characteristic of cannabis flowers	Complies	YES
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: 19.9% – 24.2%	< 0.1 % < 0.1 % 20.0 %	YES
Foreign matter	EP 3028	$\leq 2.0\%$	< 2 %	YES
Loss on drying	EP 3028	$\leq 12.0\%$	8.5 %	YES
Pesticides	EP 2.8.13	Complies with the requirements EP 2.8.13	n.n.	YES
Aflatoxins	EP 2.8.18	B1: $\leq 2 \mu\text{g/kg}$ Sum B1, B2, G1, G2: $\leq 4 \mu\text{g/kg}$	n.n. n.n.	YES
Ochratoxin A	EP 2.8.22	$\leq 20 \mu\text{g/kg}$	n.n.	
Heavy metals analysis	EP 2.4.27	Arsenic: $\leq 0.2 \text{ ppm}$ Cadmium: $\leq 0.3 \text{ ppm}$ Lead: $\leq 0.5 \text{ ppm}$ Mercury: $\leq 0.1 \text{ ppm}$	n.n. n.n. n.n. n.n.	YES
Microbiological purity		EP 5.1.8C		YES
TAMC	EP 2.6.12	$\leq 500\,000 \text{ cfu/g}$	5 000 cfu/g	
TYMC		$\leq 50\,000 \text{ cfu/g}$	18 000 cfu/g	
Bile tolerant gram neg. bacteria	EP 2.6.31	$\leq 10\,000 \text{ cfu/g}$	< 10 cfu/g	
Escherichia coli		absent / 1 g	absent / 1g	
Salmonella species		absent / 25 g	absent / 25g	

n.n. = below limit of quantitation (LOQ)

LOQ Aflatoxin B1, B2, G1: 0.5 $\mu\text{g/kg}$; LOQ Aflatoxin G2: 1 $\mu\text{g/kg}$

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

Expiry date	05/2025
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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.



03.09.2024

Dr. Klaus-Uwe Pechar

Date / Signature Qualified Person