

## Certificate of Analysis CANNABIS Flos Batch: 27402

Product	Therismos 23/1 JFG		
Country	Germany		
Manufacturing license No.:	DE_RP_01_MIA_2022_0054		
Potency	Δ <sup>9</sup> -Tetrahydrocannabinol approx. 23%		
	Cannabidiol ≤ 1%		
Dosage form	Cannabis Flowers, dried; cultivar: Jet Fuel Gelato		
Pack size	15 g metallized PET pouches		
Batch bulk	PS-PT-ARK-110923		
Batch	27402		

Grower/drier/packager	Pideka SAS, Centro Empresarial Oikos, Km 21 Via Tunja Tocancipa Cundinamarca 251010 Colombia	
Labelling	PS Pharma Service GmbH Lise-Meitner-Str. 10 40670 Meerbusch	Manufacturing Date: 15.07.2024
Manufacturing license No.:	DE_NW_03_MIA_2024_0015	

Test	Method	Specification	Result	Complies
Odour	Organoleptic	Characteristic of cannabis flowers	Complies	YES
Identification A	DAB Monograph	Complies with the description of the DAB monograph	Complies	YES
Identification B	DAB Monograph	Complies with the description of the DAB monograph	Complies	YES
Identification C	DAB Monograph (TLC) EP 2.2.27	Complies with the description of the DAB monograph	Complies	YES
Assay	DAB Monograph (HPLC) EP 2.2.29	CBD total: ≤1.0% CBN: ≤1.0% THC total: 20.7 – 25.3%	< 0.1 % < 0.1% 23.5 %	YES
Foreign matter	DAB, EP 2.8.2	≤ 2.0 %	< 2%	YES
Loss on drying	EP 2.2.32	≤ 10.0 %	9.0 %	YES
Pesticides	EP 2.8.13	Complies with the requirements EP 2.8.13	n.n.	YES
Aflatoxins	EP 2.8.18	B1: ≤ 2.0 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	Lead: ≤ 5.0 ppm Mercury: ≤ 0.1 ppm Cadmium: ≤ 1.0 ppm Arsenic: ≤ 2.0 ppm	< 0.1 ppm < 0.05 ppm 0.08 ppm < 0.1 ppm	YES
Microbiological purity		EP 5.1.8.C		YES
TAMC TYMC Bile tolerant gram neg.	EP 2.6.12 EP 2.6.31	≤ 500 000 cfu/g ≤ 50 000 cfu/g	< 10 000 cfu/g 21 000 cfu/g	
bacteria Esherichia coli Salmonella species		≤ 10 000 cfu/g absent /1g absent /25g	< 10 000 cfu/g absent /g absent /25g	

n.n. = below limit of quantitation

Examined by:

QSI GmbH, Flughafendamm 9a, 28199 Bremen

12/2024 Expiry date

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

Steuer Nummer 22/650/01532 VAT Nummer DE 815 754 017

T +49 261 450 98 20 F +49 261 450 98 210 info@adrexpharma.com www.adrexpharma.com Gerichtsstand: Koblenz

Sparkasse Koblenz IBAN: DE93 5705 0120 0000 2710 80 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.

M. k.

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

16.07.2024

\_Dr. Klaus-Uwe Pechar

Date / Signature Qualified Person