

neuraxpharm Arzneimittel GmbH

Certificate of Analysis and Release

Name, Strength/ Potency	NAXIVA-PANAXOL CBD 25 / 30 ml Cann. Ext.
Dosage Form	standardized Cannabis extract (CBD 2.6% w/w; 25 mg/ml) in MCT
Batch Number	22/206-MT
Manufacturing Date	23.01.2023
Expiry Date	12/2024

Test	Specification	Results
Organoleptic properties (visual Inspection)	Homogenous greenish or yellow to brown liquid, no significant precipitate observed	complies
Relative density (EP* 2.2.5)	Indicative for individual product	0.95 g/ml
Identification (TLC; DAB*)	Zonal pattern of test solution conforms to the zonal pattern of reference solution	complies
Assay – Cannabinoids concentration (HPLC; DAB*) CBD	95 - 105 % of CBD label claim	95.78% (2.49% (w/w))
Δ^9 THC	$\leq 6\%$ of CBD label claim	4.23% (0.11% (w/w))
Vitamin E (HPLC)	$\geq 0.4\%$ (w/w)	0.4% (w/w)
Heavy metals (EP* 2.4.27) Cadmium Lead Mercury	≤ 1.0 ppm ≤ 5.0 ppm ≤ 0.1 ppm	< 0.05 ppm < 0.1 ppm < 0.05 ppm
Toxins-Mycotoxins (EP* 2.8.18) Aflatoxin B1 Total Aflatoxins (B1, B2, G1, G2) Ochratoxin A	≤ 2 $\mu\text{g/kg}$ ≤ 4 $\mu\text{g/kg}$ ≤ 18.85 $\mu\text{g/kg}$	n.d. n.d. n.d.
Purity test (HPLC; DAB*) Cannabinol (CBN)	$\leq 2.5 \%$ (w/w)	$< 0.001\%$ (w/w)


Microbiological quality (EP* 2.6.12; 2.6.31) TAMC	10 ⁴ CFU/g (max. acceptable count: 50000 CFU/g)	< 100 CFU/g
TYMC	10 ² CFU/g (max. acceptable count: 500 CFU/g)	< 10 CFU/g
<i>Bile-tolerant gram-negative bacteria</i> <i>Escherichia coli</i> <i>Salmonella</i>	<10 ² CFU/g Absence (1 g) Absence (25 g)	< 10 CFU/g negative negative
Residual Solvents Ethanol (GC) Water content (EP* 2.5.32)	≤ 5000 ppm ≤ 0.5 % (w/w)	n.d. < 0.2%
Pesticides (EP*2.8.13)	Complies with limits indicated in EP*2.8.13	complies

*current edition

Abbreviations: n.d.: not detectable; ppm: parts per million; EP: european pharmacopoeia; TLC: thin layer chromatography; TAMC: total aerobic microbial count; TYMC: total combined yeasts and molds count; CFU: colony-forming unit.

This batch of product has been manufactured and tested in full compliance with the EU-GMP requirements. The tested batch complies with the specification and can be released.

28. APR. 2023



Issued by:
M. Stiwinski
Quality Assurance

28. APR. 2023



Approved by:
Dr. Claudia Scholzen
Deputy Head of QC

16. MAI 2023



Approved by:
Dr. Jean-Paul M. Makosi
Qualified Person