

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	23/123-MT
Manufacturing Date	20.09.2023
Expiry Date	08/2025

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	0.92
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	96.92% (2.52% (w/w))
Δ^9 THC	$\leq 6\%$ of CBD label claim	3.85% (0.10% (w/w))
Vitamin E (HPLC)	$\geq 0.4\%$ (w/w)	0.5% (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; 2.8.22) 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/ml (max. acceptable count: 50000 CFU/ml)	< 10 CFU/ml
TYMC	10 ² CFU/ml (max. acceptable count: 500 CFU/ml)	< 10 CFU/ml
<i>Bile-tolerant gram-negative bacteria</i> <i>Escherichia coli</i> <i>Salmonella</i>	<10 ² CFU/ml Absence (1 ml) Absence (25 ml)	< 10 CFU/ml absent absent
Residual Solvents Ethanol (GC)	≤ 5000 ppm	n.d.
Water content (EP* 2.5.12)	≤ 0.5 %	< 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.

I hereby confirm the release of the batch by a Qualified Person of a contract manufacturer located in EU. The batch was produced and released in compliance to GMP-Guidelines and complies with the specifications.

27.03.2024


Issued by:
 M. Sliwinski
 Quality Assurance

27. MRZ. 2024


Approved by:
 Dr. M. Schroers
 Head of QC