

Product Specification Sheet

Pharma Grade CBD E-Liquid

Test Item	Method	Acceptance Criteria
Appearance	Visual inspection	Complies
CBD content	HPLC	95 -105%
THC Content	HPLC	< 0.01%
Microbiological quality	Ph. Eur. 5.1.4	Complies

• Ingredients used in our liquids are all Pharma grade

CBD used in our liquids is a high quality Pharma grade Active substance with the following specifications

Manufacturing under GMP (Good manufacturing Practice), the standard for pharmaceutical preparations

CBD Purest Active Ingredient (99.5 ± 0.5%)

Test Item	Method	Acceptance Criteria
Appearance	Visual inspection	White to slightly yellow solid material
CBD content	Ph. Eur. 2.2.29 HPLC-DAD	99.5% ± 0.5% (m/m)
Impurities limit test		
• THC	Ph. Eur. 2.2.29	< 0.01%
• THCA	HPLC-DAD	< 0.01%
Water content (w/w %)	Karl Fisher	< 0.5%
Residual solvents (w/w %)	GC	
• Ethanol		< 0.1%
• Others		< 0.05%
Aflatoxins*		
• B1:	Ph. Eur. 2.2.29	< 2µg/Kg
• Sum of B1, B2, G1 and G2	HPLC-DAD	< 4µg/Kg
Pesticides and Fungicides*	Ph. Eur. 2.8.13	All common pesticides have to be below their specific limits
Microbiological quality	Ph. Eur. 5.1.4	TAMC: ≤10 ³ CFU/g
	(2.6.12 and 2.6.13)	TYMC: ≤10² CFU/g
GMO Genetically Modified Organisms	Declaration	CBD Purest Active Ingredient is manufactured
		from not genetically modified organisms
Resin and Plant material	Declaration	CBD purest Active ingredient is from herbal
		origin, but free from resin and plant material

* measured in the hemp flowers due to matrix interferences in the manufactured product