



Certificate of Analysis

Sample: KN11103008-001

Harvest/Lot ID: 1

Batch#: C21101

Seed to Sale# N/A

Batch Date: N/A

Sample Size Received: 10 ml

Total Weight/Volume: N/A

Retail Product Size: 10 ml

ordered : 11/01/21

sampled : 11/01/21

Completed: 11/12/21

Sampling Method: SOP Client Method

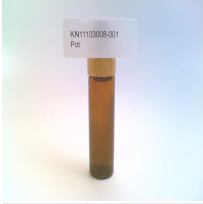










PASSED

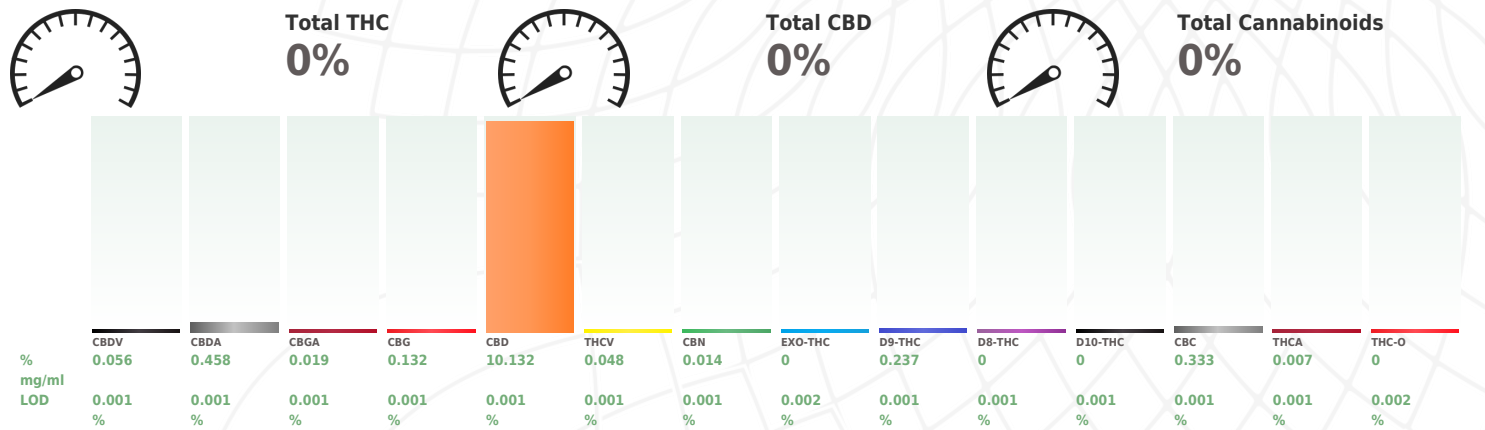
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Nov 12, 2021 | Cativa Health

1040 University Blvd
Portsmouth, VA, 23703, US



PRODUCT IMAGE	SAFETY RESULTS								MISC.
	 Pesticides NOT TESTED	 Heavy Metals NOT TESTED	 Microbials NOT TESTED	 Mycotoxins NOT TESTED	 Residuals Solvents NOT TESTED	 Filtration NOT TESTED	 Water Activity NOT TESTED	 Moisture NOT TESTED	 Terpenes NOT TESTED
	Cannabinoid								PASSED



Analyzed by: 113 **Weight:** 0.2054g **Extraction date:** 11/04/21 01:11:39 **Extracted By:** 113
Analysis Method - Expanded Measurement of Uncertainty: Flower Matrix d9-THC:12.7%, THCA: 9.5%, TOTAL THC 11.1%. These uncertainties represent an expanded uncertainty expressed at approximately the 95% confidence level using a coverage factor k=2 for a normal distribution.
Reviewed On: 11/12/21 07:50:50 **Batch Date:** 11/03/21 13:40:42 **Instrument Used:** HPLC E-SHI-008 **Running On:**
Analytical Batch: KN001507POT **Dilution:** 40
Reagent: 081321.R04; 102821.R10; 102821.R09
Consumables: 947B9291.217; 0030220

Full spectrum cannabinoid analysis utilizing High Performance Liquid Chromatography with UV/PDA detection (HPLC-UV/PDA). (Method: SOP.T.30.031.TN for sample prep and Shimadzu High Sensitivity Method SOP.T.40.020 for analysis.) *Based on FL action limits.

This report shall not be reproduced, unless in its entirety, without written approval from Kaycha Labs. This report is an Kaycha Labs certification. The results relate only to the material or product analyzed. Test results are confidential unless explicitly waived otherwise. Void after 1 year from test end date. Cannabinoid content of batch material may vary depending on sampling error. IC=In-control QC parameter, NC=Non-controlled QC parameter, ND=Not Detected, NA=Not Analyzed, ppm=Parts Per Million, ppb=Parts Per Billion. Limit of Detection (LoD) and Limit of Quantitation (LoQ) are terms used to describe the smallest concentration that can be reliably measured by an analytical procedure. RPD=Reproducibility of two measurements. Action Levels are State determined thresholds for human safety for consumption and/or inhalation. The result >99% are variable based on uncertainty of measurement (UM) for the analyte. The UM error is available from the lab upon request. The "Decision Rule" for the pass/fail does not include the UM. The limits are based on F.S. Rule 64-4.310.

Sue Ferguson

Lab Director

State License # n/a
ISO Accreditation # 17025:2017


Signature

11/12/21

Signed On