



# Certificate of Analysis

Sample:KN20412004-003  
Harvest/Lot ID: 002  
Batch#: 002  
Seed to Sale# N/A  
Batch Date: N/A  
Sample Size Received: 10 ml  
Total Weight/Volume: N/A  
Retail Product Size: 10 units  
ordered : 04/08/22  
sampled : 04/08/22  
Completed: 04/14/22 Expires: 04/14/23  
Sampling Method: SOP Client Method

Apr 14, 2022 | Cativa Health  
1040 University Blvd  
Portsmouth, VA, 23703, US



**PASSED**  
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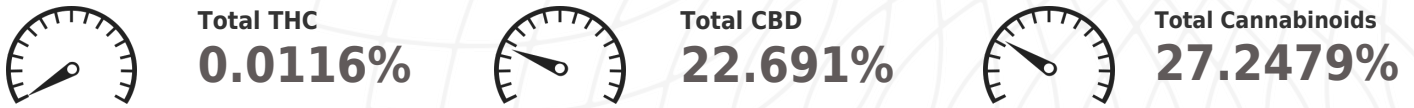
PRODUCT IMAGE SAFETY RESULTS



 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
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MISC.

 **Cannabinoid** **PASSED**



	TOTAL THC	TOTAL CBD	TOTAL CBG	CBDV	CBDA	CBGA	CBG	CBD	THCV	CBN	EXO-THC	D9-THC	D8-THC	D10-THC	CBC	THCA	D8-THCO	D9-THCO	THC-O
%	0.0116	22.691	4.4145	0.1172	ND	<0.01	4.4145	22.691	ND	<0.01	ND	0.0116	ND	ND	0.0136	<0.01	ND	ND	ND
mg/ml	0.1113	217.8336	42.3792	1.1251	ND	<0.096	42.3792	217.8336	ND	<0.096	ND	0.1113	ND	ND	0.1305	<0.096	ND	ND	ND
LOD	0.001	0.001	0.001	0.001	0.001	0.001	0.001	0.001	0.001	0.001	0.002	0.001	0.001	0.001	0.001	0.001	0.002	0.002	0.002
%	%	%	%	%	%	%	%	%	%	%	%	%	%	%	%	%	%	%	%

Cannabinoid Profile Test

Analyzed by: 113	Weight: 0.2069g	Extraction date: 04/14/22 09:04:39	Extracted By: 113
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Dilution: 40  
Reagent: 081321.R04, 041122.R08, 040522.R04  
Consumables: 947.251, 12123-046CC-046  
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.40.031 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

04/14/22

Signed On