



# Certificate of Analysis

Feb 21, 2020 | Formulated Wellness

5021 s state rd 7 davie  
Florida, United States 33314



Sample: DA00219010-001

Harvest/Lot ID: N/A

Seed to Sale #N/A

Batch Date : N/A

Batch#: 231024193

Sample Size Received: 30

Ordered : 02/18/20

Sampled : 02/18/20

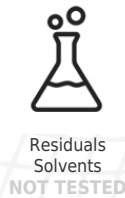
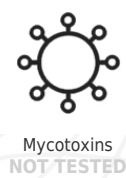
Completed: 02/21/20 Expires: 02/21/21

Sampling Method: SOP Client Method

**PASSED**

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PRODUCT IMAGE SAFETY RESULTS



MISC.

CANNABINOID RESULTS



Total THC  
**0.000%**



Total CBD  
**1.750%**



Total Cannabinoids  
**1.806%**

CBC	CBGA	CBG	THCV	D8-THC	CBDV	CBN	CBDA	CBD	D9-THC	THCA
0.038 %	ND	ND	ND	ND	ND	0.018 %	ND	1.750 %	ND	ND
0.380 mg/g	ND	ND	ND	ND	ND	0.180 mg/g	ND	17.500 mg/g	ND	ND
0.001 ppm	0.001 ppm	0.001 ppm	0.001 ppm	0.001 ppm	0.001 ppm	0.001 ppm	0.001 ppm	0.0001 ppm	0.0001 ppm	0.001 ppm

Cannabinoid Profile Test

Analyzed by 1224	Weight 3.0802g	Extraction date : 02/19/20 12:02:22	Extracted By : 965
Analysis Method -SOP.T.40.020, SOP.T.30.050		Reviewed On - 02/21/20 09:35:49	
Analytical Batch -DA010366POT		Instrument Used : DA-LC-003 CBD	
		Batch Date : 02/19/20 11:45:25	

Reagent	Dilution	Consums. ID
021820.R02	400	76124-662
021320.R15		SFN-BX-1025
021320.R14		849C4-849AK
		840C6-840H

Full spectrum cannabinoid analysis utilizing High Performance Liquid Chromatography with UV detection (HPLC-UV). (Method: SOP.T.30.050 for sample prep and Shimadzu High Sensitivity Method SOP.T.40.020 for analysis. LOQ for all cannabinoids is 1 mg/L).

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.

**Jorge Segredo**  
Lab Director

State License # n/a  
ISO Accreditation # 97164



Signature

02/21/2020

Signed On