

CA-Monrovia, CA, 91016, US

Certificate of Analysis

or Analys	IS			Complet	Orde	oduct Size: 30 ered : 09/15/20 pled : 09/15/20 cpires: 09/18/21
Sep 18, 2020 Pharm	aCanna		7			SSED
Wellington, FL, 33414		Pr	arma Canna		Pag	e 1 of 2
PRODUCT IMAGE SAFETY RESULTS						MISC.
Pesticides NOT TESTED CANNABINOID RESULTS			duals ents ESTED	Water Activity NOT TESTED	Moisture	Terpenes NOT TESTED
Total THC 0.172%		Total CBE	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~		.965%	
CBDV CBD CBG THCV CBDA 0.024% 4.533% 0.086% <loq< td=""> 0.030% 0.240 45.330 0.860 ND 0.300 mg/g mg/g mg/g ND mg/g LOD 0.0001 0.0001 0.0001 0.0001 % % % % %</loq<>	CBGA CBN D9-THI 0.120% <loq< td=""> 0.172% 1.200 mg/g 1.720 mg/g ND mg/g 0.0001 0.0001 0.0001 % % %</loq<>	6 <loq <loq<br="">ND ND</loq>	THCA-A <loq ND 0.0001 %</loq 			
1068 0.514g NA Analysis Method -SOP.T.40.020, SOP.T.30.050	traction date : .C-2030(MO-HPLC-02) Batch Dat	Extracted By : NA te : 09/16/20 09:57:28				

Dilution Reagent

Consums. ID

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 0.5 mg/L). The results are reported on a dry weight basis.

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 Detectod, NA=Not Analyzed, pm=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.

Haifei Yin Lab Director

State License # NA ISO Accreditation # L18-47-1

Signature

09/18/2020

Signed On

Kaycha Labs

1500 mg N/A Matrix: Derivative

Sample:CA00915001-003

Sample Size Received: 30 gram

Harvest/Lot ID: N/A Seed to Sale #n/a Batch Date :09/15/20 Batch#: SCD1519



605 E Huntington Dr #204 CA-Monrovia, CA, 91016, US Kaycha Labs

1500 mc

N/A Matrix : Derivative

PASSED

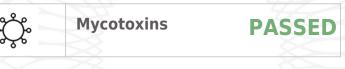
Page 2 of 2

Certificate of Analysis

PharmaCanna

2615 State Road 7, Wellington, FL, 33414 Telephone: 9543050078 Email: johnny@pharmacanna.us Sample : CA00915001-003 Harvest/LOT ID: N/A Batch# : SCD1519 Sampled : 09/15/20 Ordered : 09/15/20

Sample Size Received : 30 gram Completed : 09/18/20 Expires: 09/18/21 Sample Method : SOP Client Method



Analyte	LOD	Units	Result	Action Level (PPM)
AFLATOXIN_G2	1	ug/kg	ND	0.02
AFLATOXIN_G1	0.5	ug/kg	ND	0.02
AFLATOXIN_B2	0.5	ug/kg	ND	0.02
AFLATOXIN_B1	0.5	ug/kg	ND	0.02
OCHRATOXIN_A	5	µg/kg	ND	0.02
TOTAL AFLATOXINS (SUM OF B1, B2, G1 &G2)	4	µg/kg	ND	0.02

Analysis Method -SOP.T.30.060, SOP.T.40.060 Analytical Batch -CA000282MYC | Reviewed On - 09/17/20 11:19:19

Instrument Used : MO-LCMS-001 DER Batch Date : 09/15/20 19:06:42

Analyzed by	Weight	Extraction date	Extracted By
1051	1.011g	09/17/20 10:09:07	1051

Aflatoxins B1, B2, G1, G2, and Ochratoxins A testing using LC-MS. (Method: SOP.T.30.060 for Sample Preparation and SOP.T40.060 Procedure for Mycotoxins Quantification Using LCMS. LOQ 1.0 ppb). Total Aflatoxins (Aflotoxin B1, B2, G1, G2) must be <20µg/Kg. Ochratoxins must be <20µg/Kg.

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 Detectod, NA=Not Analyzed, pm=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.

Haifei Yin

Lab Director State License # NA ISO Accreditation # L18-47-1



Signature

09/18/2020

Signed On