



Certificate of Analysis

Sample: CA00915001-002

Harvest/Lot ID: N/A

Seed to Sale #n/a

Batch Date : 09/15/20

Batch#: SCD7519

Sample Size Received: 30 gram

Retail Product Size: 30

Ordered : 09/15/20

Sampled : 09/15/20

Completed: 09/18/20 Expires: 09/18/21

Sampling Method: SOP Client Method

PASSED

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Sep 18, 2020 | PharmaCanna

2615 State Road 7,
Wellington, FL, 33414



PRODUCT IMAGE SAFETY RESULTS





Pesticides
NOT TESTED




Heavy Metals
NOT TESTED



Microbials
NOT TESTED




Mycotoxins
PASSED




Residuals Solvents
NOT TESTED



Filtration
NOT TESTED



Water Activity
NOT TESTED



Moisture
NOT TESTED

MISC.



Terpenes
NOT TESTED

CANNABINOID RESULTS



Total THC
0.084%



Total CBD
2.331%



Total Cannabinoids
2.542%

CBDV	CBD	CBG	THCV	CBDA	CBGA	CBN	D9-THC	D8-THC	CBC	THCA-A
0.009%	2.307%	0.046%	<LOQ	0.028%	0.068%	<LOQ	0.084%	<LOQ	<LOQ	<LOQ
0.090 mg/g	23.070 mg/g	0.460 mg/g	ND	0.280 mg/g	0.680 mg/g	ND	0.840 mg/g	ND	ND	ND
LOD 0.0001 %	0.0001 %	0.0001 %	0.0001 %	0.0001 %	0.0001 %	0.0001 %	0.0001 %	0.0001 %	0.0001 %	0.0001 %

Cannabinoid Profile Test

Analyzed by: 1068 Weight: 0.525g Extraction date: NA Extracted By: NA

Analysis Method - SOP.T.40.020, SOP.T.30.050
Analytical Batch - CA000284POT Instrument Used : HPLC-2030(MO-HPLC-02) Batch Date : 09/16/20 09:57:28

Reagent Dilution 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 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.

Haifei Yin
Lab Director

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



Signature

09/18/2020

Signed On



Certificate of Analysis

PASSED
PharmaCanna

 2615 State Road 7,
 Wellington, FL, 33414

Telephone: 9543050078

Email: johnny@pharmacanna.us

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Mycotoxins
PASSED

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 11:19:05

Instrument Used : MO-LCMS-001_DER

Batch Date : 09/15/20 19:06:42

Analyzed by	Weight	Extraction date	Extracted By
1051	1.009g	09/17/20 10:09:58	1051

Aflatoxins B1, B2, G1, G2, and Ochratoxins A testing using LC-MS. (Method: SOP.T.30.060 for Sample Preparation and SOP.T.40.060 Procedure for Mycotoxins Quantification Using LCMS. LOQ 1.0 ppb). Total Aflatoxins (Aflatoxin B1, B2, G1, G2) must be <20µg/Kg. Ochratoxins must be <20µg/Kg.

Haifei Yin
 Lab Director

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