



Certificate of Analysis

Sample:GA91219001-003

Harvest/Lot ID: N/A

Seed to Sale #N/A

Batch Date :N/A

Batch#: 34919

Sample Size Received: 750 mg

Total Weight/Volume: 750 mg

Retail Product Size: 750 mg gram

Ordered : 12/17/19

sampled : 12/17/19

Completed: 12/23/19

Sampling Method: SOP Client Method

PASSED

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Dec 23, 2019 | PNP Marketing Inc
d.b.a Plants Not Pills CBD

514 N Franklin St #205, Tampa, FL 33602
Tampa, Florida, 33602, USA



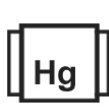
PRODUCT IMAGE



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

MISC.

CANNABINOID RESULTS



Total THC

0.000%

THC/Capsule :0.00 mg



Total CBD

3.494%

CBD/Capsule :19.87 mg



Total Cannabinoids

3.494%

Total Cannabinoids / Container
:0.000

	TOTAL CA	TOTAL CB	TOTAL TH	CBC	CBGA	CBG	THCV	DB-THC	CBDV	CBN	CBDA	CBD	THCA	D9-THC
%	3.4940	3.4940	ND	ND	ND	ND	ND	<0.010	<0.010	ND	ND	3.4940	ND	ND
mg/g	34.9400	34.9400	ND	ND	ND	ND	ND	<0.010	<0.010	ND	ND	34.9400	ND	ND
LOD	0.0000	0.0010	0.0010	0.0010	0.0010	0.0010	0.0010	0.0010	0.0010	0.0010	0.0010	0.0001	0.0010	0.0010
%	%	%	%	%	%	%	%	%	%	%	%	%	%	%

Cannabinoid Profile Test

Reagent	Dilution	Consums. ID
122019.R06	40	280678958
121919.R04		vav-09-1020
121719.R06		6970145500298
121719.R05		190715

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.

Rob Bruton
Lab Director

State License # CMTL-0001
ISO Accreditation # ISO/IEC
17025:2017 Accreditation
PJLA-Testing 97164


Signature

12/23/19

Signed On