



Certificate of Analysis

Green Lady
Rx Roll on 500mg

Sample: MO00923012-001
Harvest/Lot ID: BM IHP #2 092220
Seed to Sale #N/A
Batch Date : 09/22/20
Batch#: BM IHP #2 092220
Sample Size Received: 10 gram
Retail Product Size: 1 gram
Ordered : 09/22/20
Sampled : 09/22/20
Completed: 09/30/20 Expires: 09/30/21
Sampling Method: SOP Client Method

Sep 30, 2020 | Viobin
730 17th st
Denver, CO, 80202, USA



PASSED

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PRODUCT IMAGE SAFETY RESULTS



Pesticides
PASSED



Heavy Metals
PASSED



Microbiols
PASSED



Mycotoxins
PASSED



Residuals
Solvents
NOT TESTED



Filtration
PASSED



Water Activity
NOT TESTED



Moisture
NOT TESTED



Terpenes
NOT TESTED

MISC.

CANNABINOID RESULTS



Total THC
0.207%



Total CBD
6.551%



Total Cannabinoids
7.406%

D9-THC	THCA	CBD	CBDA	D8-THC	THCV	CBN	CBDV	CBC	CBG	CBGA
0.207%	ND	4.430%	2.419%	ND	ND	ND	ND	0.211%	0.091%	0.049%
2.070 mg/g	ND	44.300 mg/g	24.190 mg/g	ND	ND	ND	ND	2.110 mg/g	0.910 mg/g	0.490 mg/g
LOD 0.0001	0.001	0.0001	0.001	0.001	0.001	0.001	0.001	0.001	0.001	0.001
%	%	%	%	%	%	%	%	%	%	%

Filtration PASSED

Analyzed By: s Weight: NA Extraction date: NA LOD(ppm): NA Extracted By: NA

Analysis Method -SOP.T.40.013 Batch Date :
Analytical Batch -NA Reviewed On - 09/24/20 15:37:37
Instrument Used :
Running On :

This includes but is not limited to hair, insects, feces, packaging contaminants, and manufacturing waste and by-products. An SH-26/T Stereo Microscope is used for inspection.

Cannabinoid Profile Test

Analyzed by: 18 Weight: 0.2067g Extraction date: 09/23/20 02:59:58 Extracted By: 18
Analysis Method -SOP.T.40.020, SOP.T.30.050 Reviewed On - 09/24/20 09:48:44 Batch Date: 09/23/20 14:39:42
Analytical Batch -MO001127POT Instrument Used: HPLC Potency Analyzer Running On:

Reagent	Dilution	Consums. ID
	40	

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). Measurement of Uncertainty: 2.7%

This report shall not be reproduced, unless in its entirety, without written approval from Kaycha Labs. This report is on 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.

David Greene
Lab Director
State License # 19-05-02P
ISO Accreditation #
17025:2017 #97164

Signature
Signature

09/30/2020
SIGNED ON