



Certificate of Analysis

Sample: DA00806007-001
Harvest/Lot ID: G23200007
Seed to Sale #N/A
Batch Date :N/A
Batch#: G23200007
Sample Size Received: 15 ml
Retail Product Size: 15
Ordered :06/20/2022
Sampled :06/20/2022
Completed: 06/20/2022 Expires:06/20/2025
Sampling Method: SOP Client Method


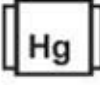

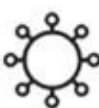





PASSED

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6/20/2022 | Medical Mary LLC
43634 Elizabeth Road Clinton Township, MI 48036

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

CANNABINOID RESULTS



Total THC
0.000%



Total CBD
0.282%



Total Cannabinoids
0.282%

CBC	CBD	CBDA	CBDV	CBG	CBGA	CBN	DB-THC	D9-THC	THCA	THCV
ND	0.282%	ND	ND	ND	ND	ND	ND	ND	ND	ND
ND	2.820 mg/lj	ND	ND	ND	ND	ND	ND	ND	ND	ND
LOD 0.001 %	0.0001 %	0.001 %	0.001 %	0.001 %	0.001 %	0.001 %	0.001 %	0.0001 %	0.001 %	0.001 %

Cannabinoid Profile Test

Analyzed by NA	Weight NA	Extraction date : NA	Extracted By : NA
Analysis Method -SOP.T.40.020, SOP.T.30.050		Reviewed On - 08/11/20 10:53:10	
Analytical Batch - Instrument Used : Batch Date :			
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 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 # CMTL-0002
ISO Accreditation # 97164



Signature

08/11/2020

Signed On