



# Certificate of Analysis

Sample: GA10105001-002  
Harvest/Lot ID: GD001  
Seed to Sale #N/A  
Batch Date : 12/30/20  
Batch#: GD001  
Sample Size Received: 1 ml  
Retail Product Size: 1  
Ordered : 12/30/20  
Sampled : 12/30/20  
Completed: 01/07/21 Expires: 01/07/22  
Sampling Method: SOP Client Method

Jan 07, 2021 | D8-Hi

2232 Dell Range Blvd.  
Cheyenne, WY, 82009, US



**PASSED**

Page 1 of 1

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
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CANNABINOID RESULTS



Total THC  
**0.000%**



Total CBD  
**0.000%**



Total Cannabinoids  
**96.233%**

CBDV	CBDA	CBGA	CBG	CBD	THCV	CBN	D9-THC	D8-THC	CBC	THCA
ND	ND	ND	ND	ND	ND	ND	ND	96.233%	ND	ND
ND	ND	ND	ND	ND	ND	ND	ND	962.330 mg/g	ND	ND
LOD 0.001%	LOD 0.001%	LOD 0.001%	LOD 0.001%	LOD 0.0001%	LOD 0.001%	LOD 0.001%	LOD 0.0001%	LOD 0.001%	LOD 0.001%	LOD 0.001%

Cannabinoid Profile Test

Analyzed by 1541	Weight 0.1047g	Extraction date : 01/05/21 04:01:36	Extracted By : 2206
Analysis Method -SOP.T.40.020, SOP.T.30.050	Reviewed On - 01/07/21 10:23:02	Batch Date : 01/05/21 15:43:57	
Analytical Batch -GA020765POT	Instrument Used : GA-HPLC-001 2030C Plus (Carl)		

Reagent	Dilution	Consums. ID
123120.R06	40	282066106
123120.R08		VAV-09-1020 Lot# 947.077
010521.06		6970145500298
		190624060
		16466-042

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.

Jeremy Campbell  
Lab Director



01/07/2021

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

Signature

Signed On