



Certificate of Analysis



CBGD0322JM
Matrix: Derivative
Accession Number: 031122UD0017
Harvest/Lot ID: CBGD0322JM
Seed to Sale: *
Batch Date: 03/08/22
Batch #: CBGD0322JM
Sample Size Received: 15 ml
Retail Product Size:
Ordered: 03/08/22
Completed: 03/15/22
COA Expires: 03/14/23
Sampling Method: SOP Client Method

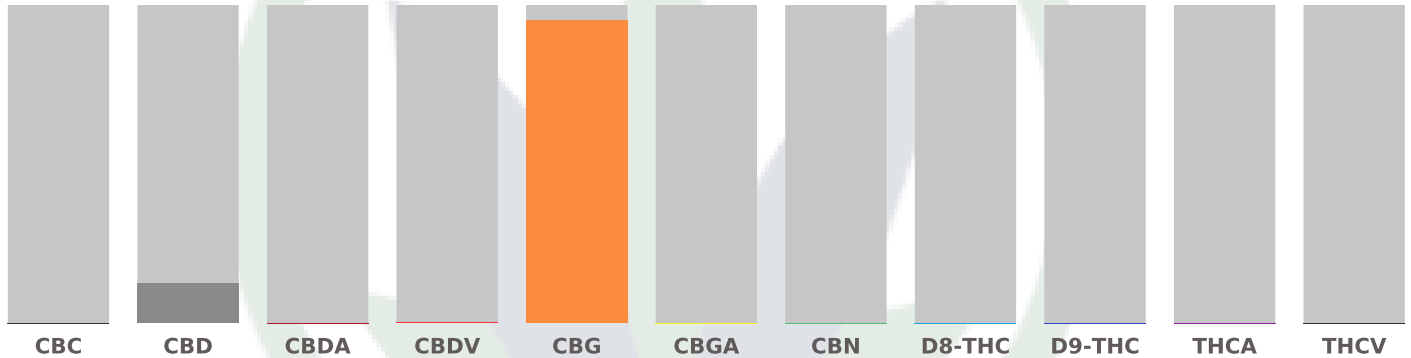
Mar 15, 2022 | Central Processors, Inc.

2413 Leaphart Rd
 West Columbia, SC, 29169
 888-654-3223



CANNABINOID RESULTS

Total THC 0.000%	Total CBD 11.203%	Total Cannabinoids 97.174%
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Conc.(wt%)	CBC	CBD	CBDA	CBDV	CBG	CBGA	CBN	D8-THC	D9-THC	THCA	THCv
ND	ND	11.203	ND	0.202	85.769	ND	ND	ND	ND	ND	ND
Conc.(mg/g)	ND	112.030	ND	2.020	857.690	ND	ND	ND	ND	ND	ND
LOQ	0.04	0.04	0.04	0.04	0.04	0.04	0.04	0.04	0.04	0.04	0.04

Analyzed by	Date	Instrument used	Analysis Method
TW	03/14/2022	Shimadzu HPLC w/ PDA	SOP.KY.02.012

Full spectrum cannabinoid analysis utilizing High Performance Liquid Chromatography with UV detection (HPLC-PDA). SOP.KY.02.005 for sample prep and SOP.KY.02.012 for analysis. % = %w/w = Percent (Weight of Analyte/Weight Product) Total Cannabinoids result reflects the absolute sum of all cannabinoids detected. **Total Potential THC/CBD is calculated using the following formulas to take into account the loss of a carboxyl group during decarboxylation Total THC = THC + (THCa*0.877) Total CBD = CBD + (CBDa*0.877)

This report shall not be reproduced, unless in its entirety, without written approval from BlueLeaf Laboratory. This report is an BlueLeaf Laboratory 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.

Daniel Burriss

Lab Director
 State License # 19-05-02P
 ISO/IEC 17025:2017

03/15/22



Signature

Signed On