



Certificate of Analysis



Simply Berry Caffeine
Matrix: Derivative
Accession Number: 062921UD0009
Harvest/Lot ID:
Seed to Sale: *
Batch Date: 06/29/21
Batch #:
Sample Size Received: 74 ml
Retail Product Size: 74
Ordered: 06/29/21
Completed: 07/22/21
Expires: 07/02/22
Sampling Method: SOP Client Method

Jul 22, 2021 | Hectare's CBD
Innovations

Louisville, KY,
(502) 548-3148

CANNABINOID RESULTS

Total THC 0.000% THC/Container :0 mg	Total CBD 0.041% CBD/Container :30.34 mg	Total Cannabinoids 0.041% Cannabinoids/Container :30.34 mg
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	CBC	CBD	CBDA	CBDV	CBG	CBGA	CBN	D8-THC	D9-THC	THCA	THCV
Conc.(wt%)	ND	0.041	ND	ND	ND	ND	ND	ND	ND	ND	ND
Conc.(mg/g)	ND	0.410	ND	ND	ND	ND	ND	ND	ND	ND	ND
LOQ	0.001	0.0001	0.001	0.001	0.001	0.001	0.001	0.001	0.0001	0.001	0.001

Analyzed by	Date	Instrument used	Analysis Method
TW	06/30/2021	Shimadzu HPLC w/ PDA	

Full spectrum cannabinoid analysis utilizing High Performance Liquid Chromatography with UV detection (HPLC-PDA). (Method: SOP.KY.02.005) sample prep and Shimadzu High Sensitivity Method SOP.KY.02.012 for analysis. LOQ for all cannabinoids is 1 mg/L. % = %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)

Filth & Foreign Matter	PASSED
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Analyzed by	Date	Instrument used	Analysis Method
DB	07/02/2021	Microscope (Amscope)	

This includes but is not limited to hair, insects, feces, packaging contaminants, and manufacturing waste and by-products. An SH-2B/T Stereo Microscope is use for inspection. SOP.KY.02.11

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

07/22/21

Signature

Signed On



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Hectare's CBD Innovations



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Table with 12 columns: Pesticides, LLOQ, Result, Units, Action Level, Pass / Fail, Pesticides, LLOQ, Result, Units, Action Level, Pass / Fail. Includes a large 'PASSED' watermark and lists various pesticides like ABAMECTIN B1A, ACEPHATE, etc.

Summary table with 4 columns: Analyzed by (DB), Date (07/02/2021), Instrument used (Shimadzu LCMSMS 8060), Analysis Method.

Pesticide screening is performed using LC/MS/MS which can screen down to below single digit ppb concentrations for the 57 pesticides analyzed. (Method: SOP.KY.02.022)

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Daniel Burriss, Lab Director, Signature, 07/22/21, Signed On, State License # 19-05-02P, Page 2 of 3



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Mycotoxins						PASSED					
Analyte	LLOQ	Result	Units	Action Level	Pass / Fail	Analyte	LLOQ	Result	Units	Action Level	Pass / Fail
Aflatoxin B1	0.001	ND	ppm	0.2	PASS	Aflatoxin B2	0.001	ND	ppm	0.2	PASS
Aflatoxin G1	0.001	ND	ppm	0.2	PASS	Aflatoxin G2	0.001	ND	ppm	0.2	PASS
Ochratoxin A+	0.001	ND	ppm	0.2	PASS						

Aflatoxins B1, B2, G1, G2, and Ochratoxins A testing using LC/MS/MS. (Method: SOP.KY.02.022)

Residual Solvents						PASSED					
Solvent	LLOQ	Result	Units	Action Level (PPM)	Pass/Fail						
2-Propanol	60.0	ND	ppm	5000	PASS						
Acetone	60	ND	ppm	5000	PASS						
Acetonitrile	60	ND	ppm	410	PASS						
Butane	200	ND	ppm	5000	PASS						
Ethyl Acetate	60	ND	ppm	5000	PASS						
Ethyl Ether	40	ND	ppm	5000	PASS						
Heptane	40	ND	ppm	5000	PASS						
Hexane	40	ND	ppm	290	PASS						
Isobutane	200	ND	ppm	5000	PASS						
M/P-Xylene	80	ND	ppm	2170	PASS						
Methanol	40	ND	ppm	3000	PASS						
O-Xylene	40	ND	ppm	2170	PASS						
Pentane	60	ND	ppm	5000	PASS						
Propane	400	ND	ppm	5000	PASS						
Toluene	40	ND	ppm	890	PASS						
Total Xylenes	120	ND	ppm	2170	PASS						

Analyzed by	Date	Instrument used	Analysis Method
DB	07/01/2021	Shimadzu GC 2010+	

Residual solvents testing for 16 common extraction solvents is performed via GC/MS. (Method: SOP.KY.02.024)

Heavy Metals						PASSED					
Metal	LLOQ	Result	Unit	Action Level	Pass / Fail						
Arsenic		ND	ppm	3	PASS						
Cadmium		ND	ppm	0.3	PASS						
Lead		ND	ppm	10	PASS						
Mercury		ND	ppm	3	PASS						

Analyzed by	Date	Instrument used	Analysis Method
DB	07/01/2021	Shimadzu ICP/MS	

Heavy Metals screening is performed using ICP-MS (Inductively Coupled Plasma - Mass Spectrometer) which can screen for toxic heavy metals (Arsenic, Cadmium, Lead, and Mercury). (Method SOP.KY.02.020)

Microbials		PASSED	
Analyte	Result		
Aspergillus Flavus	not present in 1 gram.		
Aspergillus Fumigatus	not present in 1 gram.		
Aspergillus Niger	not present in 1 gram.		
Aspergillus Terreus	not present in 1 gram.		
E. Coli	not present in 1 gram.		
Salmonella	not present in 1 gram.		

Analyzed by	Date	Instrument used	Analysis Method
DG	06/30/2021	PathogenDX	

Microbiological testing for Fungal and Bacterial Identification via Polymerase Chain Reaction (PCR) method consisting of sample DNA amplified via tandem Polymerase Chain Reaction (PCR) as a crude lysate which avoids purification. (Method SOP.KY.02.018) If a pathogenic Escherichia Coli, Salmonella, Aspergillus fumigatus, Aspergillus flavus, Aspergillus niger, or Aspergillus terreus is detected in 1g of a sample, the sample fails the microbiological-impurity testing.

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