BlueLeaf Laboratory 673 N. Bardstown Rd. Mount Washington, KY, 40047 (502) 444-2044 www.blueleaflaboratory.com Lic # 19-05-02P



Certificate of Analysis

Mar 02,2022 | Botanical Processing LLC

Louisville, Kentucky, (502) 742-7151





220215

Matrix: Topical

Accession Number: 022422UD0006

Harvest/Lot ID: Seed to Sale: *

Batch Date: 02/24/22

Batch #:

Sample Size Received: 1 units

Retail Product Size:

Ordered: 02/24/22 **Completed:** 03/02/22

Expires: 03/01/23

Sampling Method: SOP Client Method

CANNABINOID RESULTS

Total THC	Total CBD	Total Cannabinoids
0.054%	3.723%	4.046%



Analyzed by Date Instrument used Analysis Method
TW 03/01/2022 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. Voli 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 commption 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



Dang Bris

03/02/22

Signature

Signed On

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Certificate of Analysis

Botanical Processing LLC

Louisville, Kentucky,

Telephone: (502) 742-7151 Email: customercare@botanical-

0.001





220215

Matrix: Topical

Accession Number: 032122UD0026

Harvest/Lot ID:

Seed to Sale: *

Batch Date: 03/21/22

Batch #:

Sample Size Received: 1 units

Retail Product Size:

Ordered: 03/21/22 Completed: 03/25/22

COA Expires: 03/24/23

Sampling Method: SOP Client Method

MV	coto	xins
/		71

Aflatoxin G1

PASSE **Pass** Pass / Action **Analyte LLOQ Units Analyte LLOQ** Result **Units** Result Fail Fail Level Level Aflatoxin B1 0.001 ND ppm PASS Aflatoxin B2 0.001 ND ppm PASS

PASS

PASS

Aflatoxin G2

Analysis Method Analyzed by Date Instrument used DB 03/23/2022 Shimadzu LCMSMS 8060 SOP.KY.02.022

ppm

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

PASSED Heavy Metals

ND

ND

Metal	LLOQ	Result	Unit	Action Level	Pass / Fail
Arsenic	0.2	ND	ppm	2	PASS
Cadmium	0.2	ND	ppm	2	PASS
Lead	0.2	ND	ppm	5	PASS
Mercury	0.2	ND	ppm	1	PASS

Instrument used **Analysis Method** Analyzed by Date Shimadzu ICP/MS SOP.KY.02.020

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

PASSE

Analyte

Aspergillus Flavus Aspergillus Fumigatus Aspergillus Niger

Aspergillus Terreus

E. Coli

Salmonella

Analyzed by 03/24/2022

0.001

ND

Instrument used PathogenDX

Analysis Method SOP.KY.02.018

Result

not present in 1 gram.

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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03/25/22

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