

540 E Vilas Rd Suite F Central Point, OR, 97502, US

Certificate of Analysis

Kaycha Labs

CBD Support Inflamma N/A

Sample Type: Tincture

Sample: CE20427003-002 Harvest/Lot ID: INF22801E

Batch#: INF22801E

Metrc Source Package #: N/A

Metrc #: N/A

Batch Date: N/A Sample Size Received: 10 gram

> Total Weight/Volume: N/A Retail Product Size: N/A gram

> > ordered: 04/27/22 sampled: 04/27/22

Completed: 04/29/22 Sampling Method: SOP-024

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Apr 29, 2022 | Indomira/Green Earth Medicinals

License # R&D 2305 Ashland St, Ste C360 Ashland, OR, 97520, US



PRODUCT IMAGE

SAFETY RESULTS









Microbials





Residuals Solvents





Water Activity







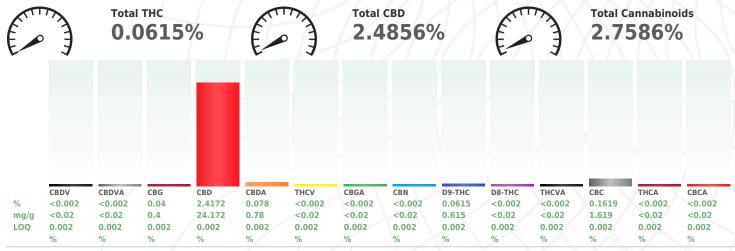


MISC.

NOT TESTED

Cannabinoid





Cannabinoid Profile Test

Analytical Batch -CE001042POT

Analyzed by 540.14.11.12

Weight 1.009g Analysis Method -SOP.T.40.020, SOP.T.30.050

Extraction date: 04/28/22 14:30:33

Reviewed On - 04/29/22 12:06:15

Instrument Used: HPLC 2030 EID 005 - Low Concentration

Extracted By:

Batch Date: 04/28/22 14:26:09

Dilution: 800

Reagent: 032922.R01: 040822.04: 120920.02

Consumables: 21/07/20; 210407; 031022-A; ASC000G11324BSF; 12315-120CC-120D; 933C4-933AL; 00321166--6 00280879 00321305-4 00321165-6 00322250-6; 2132 81421

"Total THC" and "Total CBD" are calculated values and are an Oregon reporting requirement (OAR 333-064-0100). For Cannabinoid analysis, only delta 9-THC, THCA, CBD, CBDA are ORELAP accredited analytes. Cannabinoid values reported for plant matter are dry weight corrected; Instrument LOQ for all cannabinoids is 0.5 ug/mL, LOQ is reported 'in matrix' and dependent on extraction parameters. FD = Field Duplicate; LOQ = Limit of Quantitation.

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Anthony Smith

State License # 010-10166277B9D ISO Accreditation # 99861

Signature

04/29/22

Signed On



Central Point, OR, 97502, US

Kaycha Labs

CBD Support Inflamma

N/A Sample Type : Tincture



POTENCY BATCH QC REPORT

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METHOD BLANK

Cannabinoid	LOQ	Result	Units
D9-THC_WET	0.002	0	%
THCA_WET	0.002	0	%
CBD_WET	0.002	0	%
CBDA_WET	0.002	0	%
CBN_WET	0.002	0	%
CBDV_WET	0.002	0	%
D8-THC_WET	0.002	0	%
THCV_WET	0.002	0	%
CBG_WET	0.002	0	%
CBGA_WET	0.002	0	%
CBC_WET	0.002	0	%
CBDVA_WET	0.002	0	%
THCVA_WET	0.002	0	%
CBC-A_WET	0.002	0	%

Analytical Batch - CE001042POT

Instrument Used: HPLC 2030 EID 005 - Low Concentration



LCS

Cannabinoid	LOQ	Recovery	Units	Recovery Limits
CBG_WET	0.002	95.1	%	80-120
CBD_WET	0.002	100.1	%	90-110
CBDA_WET	0.002	100	%	90-110
CBGA_WET	0.002	100.5	%	80-120
CBN_WET	0.002	100.4	%	80-120
D9-THC_WET	0.002	100.5	%	90-110
D8-THC_WET	0.002	108.5	%	90-110
CBC_WET	0.002	100	%	80-120
THCA_WET	0.002	100.5	%	90-110
CBC-A_WET	0.002	99.8	%	80-120

Analytical Batch - CE001042POT

Instrument Used: HPLC 2030 EID 005 - Low Concentration

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Anthony Smith

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Signature

04/29/22

Signed On

KML Laboratories, Inc.



261 Great Northern Road Bonners Ferry, Idaho 83805 Phone: 208-267-0818 Fax: 208-267-0878

Email: Info@kmlmicro.com

Certificate of Analysis

Green Earth Medicinals 2305 Ashland St, Suite C360 Ashland, OR 97520

Phone: 888-620-1110

Fax:

Invoice Number: 22.0559 PO Number: 04/27/22 Received Date: 04/29/2022 Number of Samples: 01

Project Name: Routine Testing

Microbiology Report:

Lab #: 22-4055	Sample Lot: INF-22801E	Sample Date: 04/27/2022
Sample Name: CBD + Inflamma Support	Additional ID:	Plated Date: 04/29/2022
Qualifying Material Number: No QM		12.7

Test Performed	Results	Units	Detection Limit	Method	Date Analyzed
Aerobic Plate Count	100	cfu/g	10	USP 43-NF 38 <2021>	05/02/2022
Coliforms	nd	cfu/g	10	Bam C4 sec G	04/30/2022
E. coli	absent	P/A	1	USP 43-NF 38 <2022>	05/02/2022
Staph aureus	absent	P/A	1	USP 43-NF 38 <2022>	05/02/2022
Yeast	nd	cfu/g	10	USP 43-NF 38 <2021>	05/04/2022
Mold	nd	cfu/g	10	USP 43-NF 38 <2021>	05/04/2022
Salmonella	absent	P/A	1	USP 43-NF 38 <2022>	05/03/2022
Pseudo. aeruginosa	absent	P/A	1	USP 43-NF 38 <m62></m62>	05/02/2022

300 Mering Companies with Res Approved By: QA Director SMV 05/04/2022



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AT-1805

Note: On this date, this material met the specifications designated above, and is not known if statistically representative of the lot evaluated on a routine basis. This information is not intended to relieve the purchaser from its responsibility to determine the suitability of this material for purchaser's purposes, to comply with all laws and regulations regarding the safe use of this material. nd = none detected above the listed detection limit

KML Laboratories, Inc.



261 Great Northern Road Bonners Ferry, Idaho 83805 Phone: 208-267-0818 Fax: 208-267-0878

Email: Info@kmlmicro.com

Certificate of Analysis

Green Earth Medicinals 2305 Ashland St, Suite C360 Ashland, OR 97520

Phone: 888-620-1110

Fax:

Invoice Number: 22.0559 PO Number: 04/27/22 Received Date: 04/29/2022 Number of Samples: 01

Project Name: **Routine Testing**

Microbiology Report:

Lab #: Control 04292022	Additional ID: Negative Control Purposes	Plated Date: 04/29/2022
Sample Name: Control 04292022		
O 116 1 March 1 N 1 OM 06 001		

Qualifying Material Number: QM-06-001

Test Performed	Results	Units	Detection Limit	Method	Date Analyzed
Aerobic Plate Count	nd	cfu/ml	10	USP 43-NF 38 <2021>	05/02/2022
Coliforms	nd	cfu/ml	10	Bam C4 sec G	04/30/2022
E. coli	absent	P/A	1	USP 43-NF 38 <2022>	05/02/2022
Staph aureus	absent	P/A	1	USP 43-NF 38 <2022>	05/02/2022
Yeast	nd	cfu/ml	10	USP 43-NF 38 <2021>	05/04/2022
Mold	nd	cfu/ml	10	USP 43-NF 38 <2021>	05/04/2022
Salmonella	absent	P/A	1	USP 43-NF 38 <2022>	05/03/2022
Pseudo. aeruginosa	absent	P/A	1	USP 43-NF 38 <m62></m62>	05/02/2022

200 Mering Companies with Approved By: QA Director SMV 05/04/202



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