

540 E Vilas Rd Suite F Central Point, OR, 97502, US (541) 668-7444

Kaycha Labs

CBD Oral Sublingual Drops Cinnamon

Matrix: Infused Product Type: Tincture

Sample:CE30502001-001

Batch#: RDCC-239017

Sample Size Received: 10 gram

Ordered: 05/02/23 Sampled: 05/02/23 Completed: 05/04/23

Revision Date: 05/30/23

Sampling Method: SOP.T.20.010.OR; ORELAP SOP-001 & -002; or Client Sampled

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Certificate of Analysis

May 30, 2023 | Indomira/Green Earth Medicinals

License # AG-R1084850IHH

2305 Ashland St, Ste C360 Ashland, OR, 97520, US

PRODUCT IMAGE

SAFFTY RESULTS















AINDOMIRA

Green Earth Medicinal

Residuals Solvents









NOT TES



MISC.

TESTED



Cannabinoid

Total THC

0.0531%



2.1074%



Total Cannabinoids 2.3093%



Analysis Method: N/A Analytical Batch: CE002500POT Instrument Used: HPLC 2030 EID 0055

Reviewed On: 05/30/23 14:23:06 Batch Date: 05/03/23 09:40:35

Analyzed Date : N/A

Reagent: 030123.R01; 031023.01

Consumables: 21/12/28: 080922-C: 210411: ASC000H02026BSF: 12543-225CD-225C: 041C-041AL: 046C6-046H: 00331867-5 00333720-5 00332100-2 00331868-5

Pipette: Fisherbrand Elite 100-1000ul EID: 0180; VWR 100-1000ul EID: 0181

"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, delta-8-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, ND= Not Detected

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Stephanie Moon

Lab Directo

State License # 010-10166277B9D ISO 17025 Accreditation # 99861



Revision: #1 - potency retest per Revision: #2 - potency retest per

Signature 05/04/23



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CBD Oral Sublingual Drops Cinnamon

N/A

Matrix : Infused Product Type: Tincture



POTENCY BATCH QC REPORT

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Cannabinoid	LOQ	Result	Units
CBDV	0.05	<loq< th=""><th>%</th></loq<>	%
CBDVA	0.05	<loq< th=""><th>%</th></loq<>	%
CBG	0.05	<loq< th=""><th>%</th></loq<>	%
CBD	0.05	<loq< th=""><th>%</th></loq<>	%
CBDA	0.05	<loq< th=""><th>%</th></loq<>	%
THCV	0.05	<loq< th=""><th>%</th></loq<>	%
CBGA	0.05	<loq< th=""><th>%</th></loq<>	%
CBN	0.05	<loq< th=""><th>%</th></loq<>	%
D9-THC	0.05	<loq< th=""><th>%</th></loq<>	%
D8-THC	0.05	<loq< th=""><th>%</th></loq<>	%
CBC	0.05	<loq< th=""><th>%</th></loq<>	%
THCA	0.05	<loq< th=""><th>%</th></loq<>	%
CBCA	0.05	<loq< th=""><th>%</th></loq<>	%

Sample Id - MB.CE002500POT Analytical Batch - CE002500POT Instrument Used: HPLC 2030 EID 0055

LCS

Cannabinoid	LOQ	Recovery	Units	Recovery Limits
CBG	0.05	94.7	%	85-115
CBD	0.05	93.1	%	90-110
CBDA	0.05	93.8	%	90-110
CBGA	0.05	96.9	%	85-115
CBN	0.05	93.6	%	85-115
D9-THC	0.05	94.4	%	90-110
D8-THC	0.05	97.5	%	90-110
CBC	0.05	94.5	%	85-115
THCA	0.05	98.3	%	90-110

Sample Id - LCS.CE002500POT Analytical Batch - CE002500POT Instrument Used: HPLC 2030 EID 0055

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Stephanie Moon

Lab Director

State License # 010-10166277B9D ISO 17025 Accreditation # 99861

Signature 05/04/23

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: 23.0576

PO Number: 050123

Received Date: 05/03/2023 Number of Samples: 01

Project Name: Routine Testing

Microbiology Report:

Lab #: 23-3695	Sample Lot: RDCC2-239017	Sample Date: 05/01/2023
Sample Name: CBD Oral Sublingual Drops Cinnamon	Additional ID: RDCC-239017	Plated Date: 05/03/2023
Qualifying Material Number: No OM		

Test Performed	Results	Units	Detection Limit	Method	Date Analyzed
				7	
Aerobic Plate Count	<10	cfu/ml	10	USP 43-NF 38 <2021>	05/06/2023
Coliforms ‡	<10	cfu/ml	10	02-228	05/04/2023
E. coli	absent	P/A	1	USP 43-NF 38 <2022>	05/06/2023
Staph aureus	absent	P/A	1	USP 43-NF 38 <2022>	05/06/2023
Yeast	<10	cfu/ml	10	USP 43-NF 38 <2021>	05/08/2023
Mold	<10	cfu/ml	10	USP 43-NF 38 <2021>	05/08/2023
Salmonella	absent	P/A	1	USP 43-NF 38 <2022>	05/07/2023
Pseudo. aeruginosa	absent	P/A	1	USP 43-NF 38 <m62></m62>	05/06/2023

Approved By: QA Director SMV 05/08/2023

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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.



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Certificate of Analysis

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Phone: 888-620-1110

Invoice Number: 23.0576

PO Number: 050123

Received Date: 05/03/2023 Number of Samples: 01

Project Name: Routine Testing

Microbiology Report:

Lab #: Control 05032023	Additional ID: Negative Control Purposes	Plated Date: 05/03/2023
Sample Name: Control 05032023		
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>	05/06/2023
Coliforms ‡	nd	cfu/ml	10	02-228	05/04/2023
E. coli	absent	P/A	1	USP 43-NF 38 <2022>	05/06/2023
Staph aureus	absent	P/A	1	USP 43-NF 38 <2022>	05/06/2023
Yeast	nd	cfu/ml	10	USP 43-NF 38 <2021>	05/08/2023
Mold	nd	cfu/ml	10	USP 43-NF 38 <2021>	05/08/2023
Salmonella	absent	P/A	1	USP 43-NF 38 <2022>	05/07/2023
Pseudo. aeruginosa	absent	P/A	1	USP 43-NF 38 <m62></m62>	05/06/2023

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