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

Kaycha Labs

CBD Relief Roll-On FD
Matrix: Infused Product
Type: Other

Certificate of Analysis

Sample:CE30601007-002
Harvest/Lot ID: C073GL-OOF
Batch#: TRC-239016
Harvest/Batch Date: 05/31/23
Sample Size Received: 10 gram
Total Amount: 8260 gram
Retail Product Size: 10 gram
Ordered: 06/01/23
Sampled: 06/01/23
Completed: 06/05/23
Sampling Method: SOP.T.20.010.OR; ORELAP SOP-001 & -002; or Client Sampled

Jun 05, 2023 | Indomira/Green Earth Medicinals

License # AG-R1084850IHH
2305 Ashland St, Ste C360
Ashland, OR, 97520, US

PRODUCT IMAGE

SAFETY RESULTS

Pesticides	Heavy Metals	Microbials	Mycotoxins	Residuals Solvents	Filth	Water Activity	Moisture	Homogeneity Testing
NOT TESTED	NOT TESTED	NOT TESTED	NOT TESTED	NOT TESTED	NOT TESTED	NOT TESTED	NOT TESTED	NOT TESTED

MISC.

Terpenes
NOT TESTED

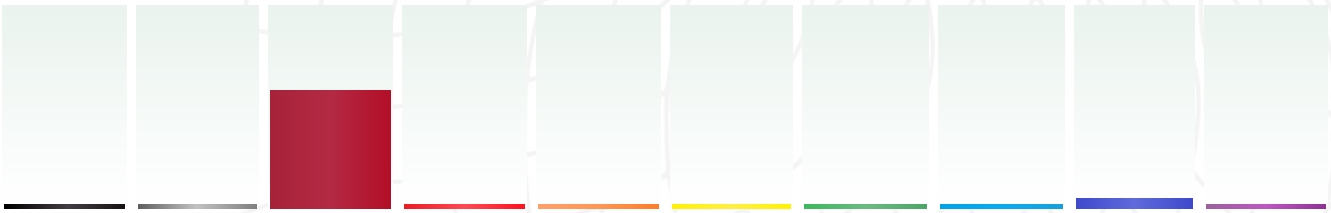
Cannabinoid

PASSED

Total THC
<LOQ

Total CBD
0.5942%

Total Cannabinoids
0.6412%



	CBDV	CBG	CBD	CBDA	CBGA	CBN	D9-THC	D8-THC	CBC	THCA
%	<LOQ	<LOQ	0.5942	<LOQ	<LOQ	<LOQ	<LOQ	<LOQ	0.047	<LOQ
mg/g	<LOQ	<LOQ	5.942	<LOQ	<LOQ	<LOQ	<LOQ	<LOQ	0.47	<LOQ
LOQ	0.002 %	0.002 %	0.002 %	0.002 %	0.002 %	0.002 %	0.002 %	0.002 %	0.002 %	0.002 %

Analyzed by: 11, 12, 834
Weight: 0.824g
Extraction date: 06/02/23 10:36:45
Extracted by: 12

Analysis Method : N/A
Analytical Batch : CE002702POT
Instrument Used : HPLC 2030 EID 0055 - Low Concentration
Analyzed Date : 06/02/23 12:00:51
Reviewed On : 06/05/23 11:41:18
Batch Date : 06/02/23 10:33:29

Dilution : 820
Reagent : 042723.R12; 031023.07; 101022.05
Consumables : 22/02/21; 080922-C; ASC000H02026BSF; 12620-307CD-307D; 042C4-042AL; 00331867-5 00333720-5 00332100-2 00331868-5; 2132
Pipette : Gilson Positive Displacement 100-1000ul EID: 0152; VWR 20-200ul EID: 0182

"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

Stephanie Moon
Lab Director

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

Signature
06/05/23

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Kaycha Labs

CBD Relief Roll-On FD
N/A
Matrix : Infused Product
Type: Other

POTENCY BATCH QC REPORT

METHOD BLANK

Cannabinoid	LOQ	Result	Units
CBDV	0.002	<LOQ	%
CBDVA	0.002	<LOQ	%
CBG	0.002	<LOQ	%
CBD	0.002	<LOQ	%
CBDA	0.002	<LOQ	%
THCV	0.002	<LOQ	%
CBGA	0.002	<LOQ	%
CBN	0.002	<LOQ	%
D9-THC	0.002	<LOQ	%
D8-THC	0.002	<LOQ	%
CBC	0.002	<LOQ	%
THCA	0.002	<LOQ	%
CBCA	0.002	<LOQ	%

Sample Id - MB.CE002702POT
Analytical Batch - CE002702POT
Instrument Used : HPLC 2030 EID 0055 - Low Concentration

LCS

Cannabinoid	LOQ	Recovery	Units	Recovery Limits
CBG	0.002	97.1	%	85-115
CBD	0.002	97.2	%	90-110
CBDA	0.002	98.8	%	90-110
CBGA	0.002	97.9	%	85-115
CBN	0.002	94.8	%	85-115
D9-THC	0.002	95.6	%	90-110
D8-THC	0.002	94.9	%	90-110
CBC	0.002	97.7	%	85-115
THCA	0.002	97.2	%	90-110

Sample Id - LCS.CE002702POT
Analytical Batch - CE002702POT
Instrument Used : HPLC 2030 EID 0055 - Low Concentration

Stephanie Moon
Lab Director

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

Signature
06/05/23

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		Invoice Number:	23.0756
2305 Ashland St, Suite C360		PO Number:	060123
Ashland, OR 97520		Received Date:	06/05/2023
Phone: 888-620-1110		Number of Samples:	01
Email: Compliance@indomira.com		Project Name:	Routine Testing

Microbiology Report:

Lab #: 23-4524	Sample Lot: TRC - 239016	Sample Date: 05/31/2023
Sample Name: CBD Relief Roll-On	Additional ID:	Plated Date: 06/05/2023
Qualifying Material Number: No QM		

Test Performed	Results	Units	Detection Limit	Method	Date Analyzed
Aerobic Plate Count	nd	cfu/ml	10	USP 43-NF 38 <61>	06/08/2023
Coliforms ‡	nd	cfu/ml	10	02-228	06/06/2023
E. *coli*	absent	P/A	1	USP 43-NF 38 <62>	06/08/2023
Staph *aureus*	absent	P/A	1	USP 43-NF 38 <62>	06/08/2023
Yeast	nd	cfu/ml	10	USP 43-NF 38 <61>	06/10/2023
Mold	nd	cfu/ml	10	USP 43-NF 38 <61>	06/10/2023
Salmonella	absent	P/A	1	USP 43-NF 38 <62>	06/09/2023
Pseudo. *aeruginosa*	absent	P/A	1	USP 43-NF 38 <62>	06/08/2023

Approved By: QA Director SMV 06/12/2023

AT-1805

This report may include work not covered by KML's current ISO accreditation as indicated by ‡.

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

F10-0817 G Ref SOP 01-023

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		Invoice Number:	23.0756
2305 Ashland St, Suite C360		PO Number:	060123
Ashland, OR 97520		Received Date:	06/05/2023
Phone: 888-620-1110		Number of Samples:	01
Email: Compliance@indomira.com		Project Name:	Routine Testing

Microbiology Report:

Lab #: Control 06052023	Additional ID: Negative Control Purposes	Plated Date: 06/05/2023
Sample Name: Control 06052023		
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 <61>	06/08/2023
Coliforms ‡	nd	cfu/ml	10	02-228	06/07/2023
E. *coli*	absent	P/A	1	USP 43-NF 38 <62>	06/08/2023
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Mold	nd	cfu/ml	10	USP 43-NF 38 <61>	06/10/2023
Salmonella	absent	P/A	1	USP 43-NF 38 <62>	06/09/2023
Pseudo. *aeruginosa*	absent	P/A	1	USP 43-NF 38 <62>	06/08/2023

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