



Certificate of Analysis

Sample: CE10909001-001
Harvest/Lot ID: 6510IHH-TCC2104

Metric #: N/A

Metric Source Package #: N/A

Batch Date: 08/24/21

Batch#: TCC-21704

Sample Size Received: 1 gram

Total Weight/Volume: 78330 gram gram

Retail Product Size: 28.35 gram

Ordered : 09/09/21

sampled : 09/09/21

Completed: 09/10/21 Expires: 09/10/22

Sampling Method: SOP-024

Sep 10, 2021 | Indomira/Green Earth Medicinals

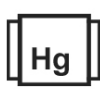
License # ODA
2305 Ashland St, Ste C360
Ashland, OR, 97520, US


Page 1 of 2

PRODUCT IMAGE



SAFETY RESULTS


Pesticides
NOT TESTED

Heavy Metals
NOT TESTED

Microbials
NOT TESTED

Mycotoxins
NOT TESTED

Residuals
Solvents
NOT TESTED

Filtration
NOT TESTED

Water Activity
NOT TESTED

Moisture
NOT TESTED

Homogeneity
NOT TESTED

Terpenes
NOT TESTED

CANNABINOID RESULTS


Total THC
0.012%

Total CBD
0.485%

Total Cannabinoids
0.536%

	CBDV	CBDVA	CBG	CBD	CBDA	THCV	CBGA	CBN	D9-THC	D8-THC	THCVA	CBC	THCA	CBCA
%	<LOQ	<LOQ	<LOQ	0.468	0.018	<LOQ	<LOQ	<LOQ	0.012	<LOQ	<LOQ	0.036	<LOQ	<LOQ
mg/g	<LOQ	<LOQ	<LOQ	4.68	0.18	<LOQ	<LOQ	<LOQ	0.12	<LOQ	<LOQ	0.36	<LOQ	<LOQ
LOQ	0.002	0.002	0.002	0.002	0.002	0.002	0.002	0.002	0.002	0.002	0.002	0.002	0.002	0.002
%	%	%	%	%	%	%	%	%	%	%	%	%	%	%

Cannabinoid Profile Test

Analyzed by 13	Weight 1.032g	Extraction date : 09/09/21 03:09:25	Extracted By : 14
Analysis Method -SOP.T.40.020, SOP.T.30.050	Reviewed On - 09/10/21 11:12:53	Batch Date : 09/09/21 15:12:12	
Analytical Batch -CE000334POT	Instrument Used : HPLC 2030 EID 005 - Low Concentration	Running On :	

Reagent	Dilution	Consums. ID	Consums. ID
090321.R01 071921.16	40	D01493069 32009E-1232 436020160AS3 436020338AS2 436021005AS3 C0000642 041CD-041C 042C4-042AL	945C6-945H F148560 0325891

"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 mg/mL, LOQ 'in matrix' is dependent on extraction parameters. FD = Field Duplicate; LOQ = Limit of Quantitation.



POTENCY BATCH QC REPORT

Page 2 of 2



METHOD BLANK

Cannabinoid	LOQ	Result	Units
CBDV_WET	0.002	<LOQ	%
CBDVA_WET	0.002	<LOQ	%
THCV_WET	0.002	<LOQ	%
CBD_WET	0.002	<LOQ	%
CBG_WET	0.002	<LOQ	%
CBDA_WET	0.002	<LOQ	%
CBN_WET	0.002	<LOQ	%
CBGA_WET	0.002	<LOQ	%
THCVA_WET	0.002	<LOQ	%
D9-THC_WET	0.002	<LOQ	%
D8-THC_WET	0.002	<LOQ	%
CBC_WET	0.002	<LOQ	%
THCA_WET	0.002	<LOQ	%
CBC-A_WET	0.002	<LOQ	%
TOTAL THC	0.002	NT	%
TOTAL CBD	0.002	NT	%
TOTAL CANNABINOIDS	0.002	NT	%
CBDV	0.002	NT	%
CBDVA	0.002	NT	%
CBG	0.002	NT	%
CBD	0.002	NT	%
CBDA	0.002	NT	%
THCV	0.002	NT	%
CBGA	0.002	NT	%
CBN	0.002	NT	%
D9-THC	0.002	NT	%
D8-THC	0.002	NT	%
THCVA	0.002	NT	%
CBC	0.002	NT	%
THCA	0.002	NT	%
CBCA	0.002	NT	%

Analytical Batch - CE000334POT

Instrument Used : HPLC 2030 EID 005 - Low Concentration



LCS

Cannabinoid	LOQ	Recovery	Units	Recovery Limits
CBG_WET	0.002	91.8	%	70-130
CBD_WET	0.002	94.2	%	70-130
CBDA_WET	0.002	95.3	%	70-130
THCV_WET	0.002	97.7	%	70-130
CBGA_WET	0.002	90.4	%	70-130
CBN_WET	0.002	92.6	%	70-130
D9-THC_WET	0.002	91.7	%	70-130
CBC_WET	0.002	92.5	%	70-130
THCA_WET	0.002	92.6	%	70-130
CBC-A_WET	0.002	93.4	%	70-130

Analytical Batch - CE000334POT

Instrument Used : HPLC 2030 EID 005 - Low Concentration



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

Kaycha Labs

Topical Relief Cream TCC-21704

N/A

Sample Type: Topical



Certificate of Analysis

Sample: CE10909001-001

Harvest/Lot ID: 65101HH-TCC2104

Metrc #: N/A

Metrc Source Package #: N/A

Batch Date: 08/24/21

Batch #: TCC-21704

Sample Size Received: 1 gram

Total Weight/Volume: 78330 gram gram

Retail Product Size: 28.35 gram

Ordered : 09/09/21

sampled : 09/09/21

Completed: 09/10/21 Expires: 09/10/22

Sampling Method: SOP-024

Sep 10, 2021 | Indomira/Green
Earth Medicinals

License # ODA
2305 Ashland St, Ste C360
Ashland, OR, 97520, US



Cannabinoid Potency Batch Quality Report:

OLCC/ODA Control Study ID: 122419-6

Sample	Lab ID	Total THC* (mg/g)	Total CBD* (mg/g)
Topical Relief Cream TCC-21704	CE10909001-001	0.12	4.85
Topical Relief Cream TCC-21704 FD	CE10909001-002	0.12	4.89
Average		0.12	4.87
RPD		0.00%	0.82%
RPD Status:		PASS	
Max Total THC [†] <		35.3 mg/g	<1000 mg†/28.35g unit
Max allowable Total THC <		38.8 mg/g	<1100 mg/28.35g unit
Highest measured		0.12 mg/g	
Average Total THC		0.12 mg/g	
Total THC per unit status		PASS	
Average Total THC per unit of sale:		3.4 mg/ unit	
Average Total CBD per unit of sale:		138.06 mg/ unit	

Analytical batch ID: 334POT



540 E. Vilas Rd., Suite F
Central Point, OR 97502
www.kaychalabs.com
541.668.7444

Anthony Smith, Ph.D

This report shall not be reproduced, unless in its entirety, without written approval from Kaycha Labs and KGO, LLC. This report is a Kaycha Labs 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. Sampling method: KGO-SOP-018; ORELAP-SOP-001, -002.

* "Total THC" and "Total CBD" are calculated values and are an Oregon reporting requirement (OAR 333-064-0100).

† OAR 333.007-0210 Retail adult use cannabis concentration and serving limits. These are *maximum* and *maximum allowable* (+10%) Total THC limits for concentration (mg/g) based on your certified CS retail unit mass, not the declared target THC from the CS.

KGO-BQR-CE10909001-001_002 Indomira-GEM

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

Invoice Number: 21.1155
PO Number: 090821
Received Date: 09/10/2021
Number of Samples: 01
Project Name: Routine Testing

Microbiology Report:

Lab #: 21-9208	Sample Lot: TCC-21704	Sample Date: 09/08/2021
Sample Name: Topical Relief Cream	Additional ID:	Plated Date: 09/10/2021
Qualifying Material Number: No QM		

Test Performed	Results	Units	Detection Limit	Method	Date Analyzed
Aerobic Plate Count	nd	cfu/g	10	USP 43-NF 38 <61>	09/13/2021
Coliforms	nd	cfu/g	10	Bam C4 sec G	09/11/2021
<i>E. coli</i>	absent	P/A	1	USP 43-NF 38 <62>	09/13/2021
<i>Staph aureus</i>	absent	P/A	1	USP 43-NF 38 <62>	09/13/2021
Yeast	nd	cfu/g	10	USP 43-NF 38 <61>	09/15/2021
Mold	nd	cfu/g	10	USP 43-NF 38 <61>	09/15/2021
Salmonella	absent	P/A	1	USP 43-NF 38 <62>	09/14/2021
<i>Pseudo. aeruginosa</i>	absent	P/A	1	USP 43-NF 38 <62>	09/13/2021

Approved By: QA Director SMV 09/15/2021

Confidential

Page 1 of 2

This Certificate/Report shall not be reproduced, except in full, without the prior written consent of KML Laboratories, Inc.

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



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

Invoice Number: 21.1155
PO Number: 090821
Received Date: 09/10/2021
Number of Samples: 01
Project Name: Routine Testing

Microbiology Report:

Lab #: Control 09102021	Additional ID: Negative Control Purposes	Plated Date: 09/10/2021
Sample Name: Control 09102021		
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>	09/13/2021
Coliforms	nd	cfu/ml	10	Bam C4 sec G	09/11/2021
<i>E. coli</i>	absent	P/A	1	USP 43-NF 38 <62>	09/13/2021
<i>Staph aureus</i>	absent	P/A	1	USP 43-NF 38 <62>	09/13/2021
Yeast	nd	cfu/ml	10	USP 43-NF 38 <61>	09/15/2021
Mold	nd	cfu/ml	10	USP 43-NF 38 <61>	09/15/2021
Salmonella	absent	P/A	1	USP 43-NF 38 <62>	09/14/2021
<i>Pseudo. aeruginosa</i>	absent	P/A	1	USP 43-NF 38 <62>	09/13/2021

Approved By: QA Director SMV 09/15/2021

Confidential

Page 2 of 2

This Certificate/Report shall not be reproduced, except in full, without the prior written consent of KML Laboratories, Inc.

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

