



Certificate of Analysis

Sample: CE10922006-001
Harvest/Lot ID: 6510IHH-INF2103

Metric #: N/A

Metric Source Package #: N/A

Batch Date: 09/21/21

Batch#: INF-21703E

Sample Size Received: 9.69 gram

Total Weight/Volume: 10915 gram gram

Retail Product Size: 1.6 gram

Ordered : 09/22/21

sampled : 09/22/21

Completed: 09/24/21 Expires: 09/24/22

Sampling Method: SOP-024

Page 1 of 2

Sep 24, 2021 | Indomira/Green
Earth Medicinals

License # ODA

2305 Ashland St, Ste C360

Ashland, OR, 97520, US



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
0.046%



Total CBD
2.358%



Total Cannabinoids
2.61%

	CBDV	CBDVA	CBG	CBD	CBDA	THCV	CBGA	CBN	D9-THC	D8-THC	THCVA	CBC	THCA	CBCA
%	<LOQ	<LOQ	0.041	2.284	0.084	<LOQ	<LOQ	<LOQ	0.046	<LOQ	<LOQ	0.153	<LOQ	<LOQ
mg/g	<LOQ	<LOQ	0.41	22.84	0.84	<LOQ	<LOQ	<LOQ	0.46	<LOQ	<LOQ	1.53	<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 11	Weight 1.03g	Extraction date : 09/23/21 04:09:23	Extracted By : 13
Analysis Method -SOP.T.40.020, SOP.T.30.050	Reviewed On - 09/24/21 11:31:32	Batch Date : 09/23/21 16:17:04	
Analytical Batch -CE000372POT	Instrument Used : HPLC 2030 EID 005 - Low Concentration	Running On :	

Reagent	Dilution	Consums. ID	Consums. ID
091721.11	40	D01493069 32009E-1232 436020160A53 436020338A52 436021005A53 C0000642 041CD-041C 042C4-042AL	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 CANNABINOIDS	0.002	<LOQ	%
TOTAL CBD	0.002	<LOQ	%
TOTAL THC	0.002	<LOQ	%
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	%
THCVA	0.002	<LOQ	%
CBC	0.002	<LOQ	%
THCA	0.002	<LOQ	%
CBCA	0.002	<LOQ	%

Analytical Batch - CE000372POT

Instrument Used : HPLC 2030 EID 005 - Low Concentration

LCS

Cannabinoid	LOQ	Recovery	Units	Recovery Limits
CBG_WET	0.002	102.5	%	70-130
CBD_WET	0.002	97.8	%	70-130
CBDA_WET	0.002	97.6	%	70-130
THCV_WET	0.002	103.1	%	70-130
CBGA_WET	0.002	95.2	%	70-130
CBN_WET	0.002	102.6	%	70-130
D9-THC_WET	0.002	101.5	%	70-130
CBC_WET	0.002	101.7	%	70-130
THCA_WET	0.002	99.1	%	70-130
CBC-A_WET	0.002	105	%	70-130

Analytical Batch - CE000372POT

Instrument Used : HPLC 2030 EID 005 - Low Concentration



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

Certificate of Analysis

Sep 24, 2021 | Indomira/Green
Earth Medicinals

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



Kaycha Labs

Inflamma Caps INF-21703E

N/A

Sample Type: Ingestible



Sample: CE10922006-001

Harvest/Lot ID: 6510IHH-INF2103

Metrc #: N/A

Metrc Source Package #: N/A

Batch Date: 09/21/21

Batch#: INF-21703E

Sample Size Received: 9.69 gram

Total Weight/Volume: 10915 gram gram

Retail Product Size: 1.6 gram

Ordered : 09/22/21

sampled : 09/22/21

Completed: 09/24/21 Expires: 09/24/22

Sampling Method: SOP-024

Cannabinoid Potency Batch Quality Report:

OLCC/ODA Control Study ID:

Sample	Lab ID	Total THC* (mg/g)	Total CBD* (mg/g)
Inflamma Caps INF-21703E	CE10922006-001	0.46	23.58
Inflamma Caps INF-21703E FD	CE10922006-002	0.47	23.43
Average		0.47	23.51
RPD		2.15%	0.64%
RPD Status:		PASS	
Max Total THC [†] <		31.3 mg/g	<50 mg†/1.6g unit
Max allowable Total THC <		34.4 mg/g	<55 mg/1.6g unit
Highest measured		0.47 mg/g	
Average Total THC		0.47 mg/g	
Total THC per unit status		PASS	
Average Total THC per unit of sale:		0.7 mg/ unit	
Average Total CBD per unit of sale:		37.61 mg/ unit	

Analytical batch ID: 372POT



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-CE10922006-001_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
Fax:

Invoice Number: 21.1220
PO Number: 092221
Received Date: 09/29/2021
Number of Samples: 01
Project Name: Routine Testing

Microbiology Report:

Lab #: 21-9778	Sample Lot: INF-21703E	Sample Date: 09/22/2021
Sample Name: Inflamm Caps	Additional ID:	Plated Date: 09/29/2021
Qualifying Material Number: No QM		

Test Performed	Results	Units	Detection Limit	Method	Date Analyzed
Aerobic Plate Count	155	cfu/g	10	USP 43-NF 38 <2021>	10/02/2021
Coliforms	nd	cfu/g	10	Bam C4 sec G	09/30/2021
<i>E. coli</i>	absent	P/A	1	USP 43-NF 38 <2022>	10/02/2021
<i>Staph aureus</i>	absent	P/A	1	USP 43-NF 38 <2022>	10/02/2021
Yeast	nd	cfu/g	10	USP 43-NF 38 <2021>	10/04/2021
Mold	nd	cfu/g	10	USP 43-NF 38 <2021>	10/04/2021
Salmonella	absent	P/A	1	USP 43-NF 38 <2022>	10/03/2021
<i>Pseudo. aeruginosa</i>	absent	P/A	1	USP 43-NF 38 <m62>	10/02/2021

Approved By: QA Director SMV 10/04/2021

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
Fax:

Invoice Number: 21.1220
PO Number: 092221
Received Date: 09/29/2021
Number of Samples: 01
Project Name: Routine Testing

Microbiology Report:

Lab #: Control 09292021	Additional ID: Negative Control Purposes	Plated Date: 09/29/2021
Sample Name: Control 09292021		
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>	10/02/2021
Coliforms	nd	cfu/ml	10	Bam C4 sec G	09/30/2021
<i>E. coli</i>	absent	P/A	1	USP 43-NF 38 <2022>	10/02/2021
<i>Staph aureus</i>	absent	P/A	1	USP 43-NF 38 <2022>	10/02/2021
Yeast	nd	cfu/ml	10	USP 43-NF 38 <2021>	10/04/2021
Mold	nd	cfu/ml	10	USP 43-NF 38 <2021>	10/04/2021
Salmonella	absent	P/A	1	USP 43-NF 38 <2022>	10/03/2021
<i>Pseudo. aeruginosa</i>	absent	P/A	1	USP 43-NF 38 <m62>	10/02/2021

Approved By: QA Director SMV 10/04/2021

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



AT-1805