



Certificate of Analysis

Sample: CE21212005-001

Harvest/Lot ID: N/A

Batch#: ASC-228023

Metric Source Package #: N/A

Metric #: N/A

Batch Date: N/A

Sample Size Received: 15 gram

Total Amount: N/A

Retail Product Size: N/A gram

Ordered: 12/12/22

Sampled: 12/12/22

Completed: 12/14/22

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

Pages 1 of 2

Dec 14, 2022 | Indomira/Green Earth Medicinals

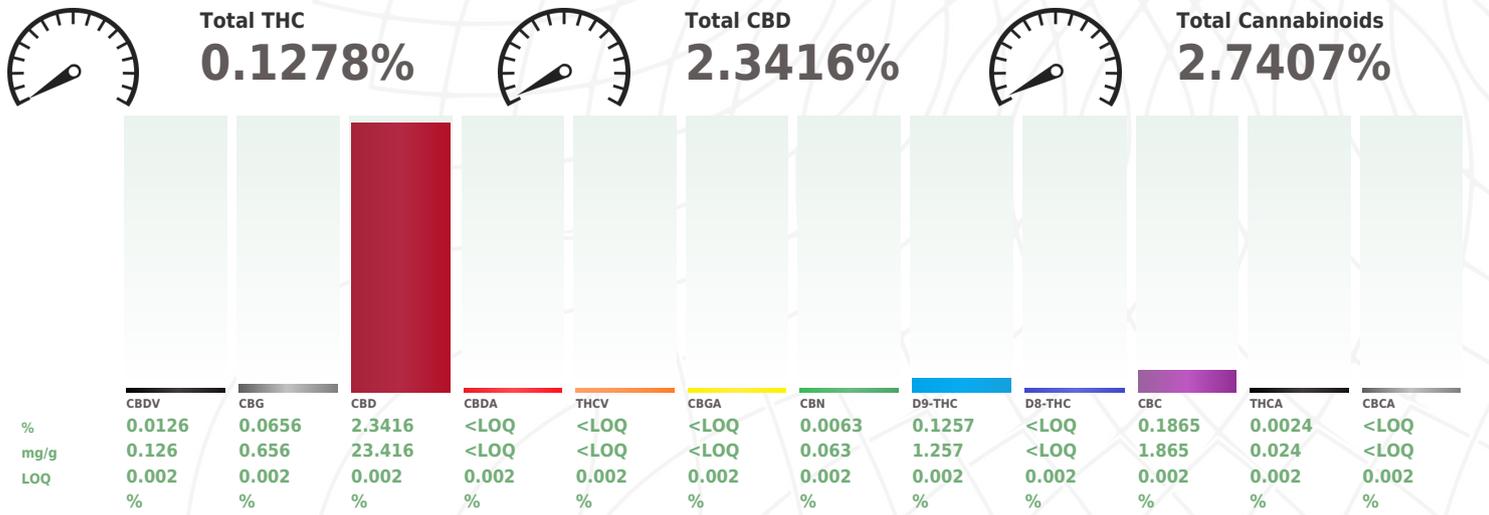
License # Indomira

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



PRODUCT IMAGE	SAFETY RESULTS								MISC.	
	 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 Testing NOT TESTED	 Terpenes NOT TESTED

 **Cannabinoid** **TESTED**



Analyzed by: 11, 12, 14 Weight: 0.9g Extraction date: 12/13/22 13:32:41 Extracted by: 14

Analysis Method: N/A Analytical Batch: CE001691POT Reviewed On: 12/14/22 11:42:24
Instrument Used: HPLC 2030 EID 0055 - Low Concentration Batch Date: 12/13/22 13:27:24
Running on: N/A

Dilution: 820
Reagent: 082522.10; 120920.02; 121322.R09; 111522.R06
Consumables: 11/21/25; 080922-C; 210411; 2210449; ASC000H02026BSF; 12543-225CD-225C; 041C-041AL; 046C6-046H; 00312590-5 0032165-6 00323608-5 282851; 2132 81421
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

This report shall not be reproduced, unless in its entirety, without written approval from Kaycha Labs. This report is an Kaycha Labs certification. Laboratory reports are for informational use only, unless indicated otherwise. 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. IC=In-control QC parameter, NC=Non-controlled QC parameter, ND=Not Detected, NA=Not Analyzed, ppm=Parts Per Million, ppb=Parts Per Billion. Limit of Detection (LoD) and Limit Of Quantitation (LoQ) are terms used to describe the smallest concentration that can be reliably measured by an analytical procedure. RPD=Reproducibility of two measurements. Action Levels are State determined thresholds variable based on uncertainty of measurement (UM) for the analyte. The UM error is available from the lab upon request. The "Decision Rule" for the pass/fail does not include the UM. The limits are based on OAR 333-007, OAR 845-025.

Anthony Smith
Lab Director

State License # 010-1016627789D
ISO 17025 Accreditation # 99861



Signature

12/14/22

Signed On



POTENCY BATCH QC REPORT

 **METHOD BLANK**

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

Analytical Batch - CE001691POT
Instrument Used : HPLC 2030 EID 0055 - Low Concentration

 **LCS**

Cannabinoid	LOQ	Recovery	Units	Recovery Limits
CBG_WET	0.002	101.2	%	80-120
CBD_WET	0.002	103.9	%	90-110
CBDA_WET	0.002	99	%	90-110
CBGA_WET	0.002	99	%	80-120
CBN_WET	0.002	99.8	%	80-120
D9-THC_WET	0.002	101.1	%	90-110
D8-THC_WET	0.002	100	%	90-110
CBC_WET	0.002	100	%	80-120
THCA_WET	0.002	94.6	%	90-110

Analytical Batch - CE001691POT
Instrument Used : HPLC 2030 EID 0055 - Low Concentration

This report shall not be reproduced, unless in its entirety, without written approval from Kaycha Labs. This report is an Kaycha Labs certification. Laboratory reports are for informational use only, unless indicated otherwise. 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. IC=In-control QC parameter, NC=Non-controlled QC parameter, ND=Not Detected, NA=Not Analyzed, ppm=Parts Per Million, ppb=Parts Per Billion. Limit of Detection (LoD) and Limit Of Quantitation (LoQ) are terms used to describe the smallest concentration that can be reliably measured by an analytical procedure. RPD=Reproducibility of two measurements. Action Levels are State determined thresholds variable based on uncertainty of measurement (UM) for the analyte. The UM error is available from the lab upon request. The "Decision Rule" for the pass/fail does not include the UM. The limits are based on OAR 333-007, OAR 845-025.

Anthony Smith
Lab Director

State License # 010-1016627789D
ISO 17025 Accreditation # 99861



Signature

12/14/22

Signed On



Certificate of Analysis

Green Earth Medicinals
 2305 Ashland St, Suite C360
 Ashland, OR 97520
 Phone: 888-620-1110
 Fax:

Invoice Number: 22.1469
 PO Number: 12/12/2022
 Received Date: 12/16/2022
 Number of Samples: 01
 Project Name: Routine Testing

Microbiology Report:

Lab #: 22-10691	Sample Lot: ASC-228023	Sample Date: 12/12/2022
Sample Name: CBD ORAL Buccal Spray	Additional ID:	Plated Date: 12/16/2022
Qualifying Material Number: No QM		

Test Performed	Results	Units	Detection Limit	Method	Date Analyzed
Aerobic Plate Count	<10	cfu/ml	10	USP 43-NF 38 <2021>	12/19/2022
Coliforms	<10	cfu/ml	10	Bam C4 sec G	12/17/2022
<i>E. coli</i>	absent	P/A	1	USP 43-NF 38 <2022>	12/20/2022
<i>Staph aureus</i>	absent	P/A	1	USP 43-NF 38 <2022>	12/20/2022
Yeast	<10	cfu/ml	10	USP 43-NF 38 <2021>	12/21/2022
Mold	<10	cfu/ml	10	USP 43-NF 38 <2021>	12/21/2022
Salmonella	absent	P/A	1	USP 43-NF 38 <2022>	12/20/2022
<i>Pseudo. aeruginosa</i>	absent	P/A	1	USP 43-NF 38 <m62>	12/20/2022

Approved By: QA Director SMV 12/21/2022

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





Certificate of Analysis

Green Earth Medicinals
 2305 Ashland St, Suite C360
 Ashland, OR 97520
 Phone: 888-620-1110
 Fax:

Invoice Number: 22.1469
 PO Number: 12/12/2022
 Received Date: 12/16/2022
 Number of Samples: 01
 Project Name: Routine Testing

Microbiology Report:

Lab #: Control 12162022	Additional ID: Negative Control Purposes	Plated Date: 12/16/2022
Sample Name: Control 12162022		
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>	12/19/2022
Coliforms	nd	cfu/ml	10	Bam C4 sec G	12/17/2022
<i>E. coli</i>	absent	P/A	1	USP 43-NF 38 <2022>	12/20/2022
<i>Staph aureus</i>	absent	P/A	1	USP 43-NF 38 <2022>	12/20/2022
Yeast	nd	cfu/ml	10	USP 43-NF 38 <2021>	12/21/2022
Mold	nd	cfu/ml	10	USP 43-NF 38 <2021>	12/21/2022
Salmonella	absent	P/A	1	USP 43-NF 38 <2022>	12/20/2022
<i>Pseudo. aeruginosa</i>	absent	P/A	1	USP 43-NF 38 <m62>	12/20/2022

Approved By: QA Director SMV 12/21/2022

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](#)

