














# Certificate of Analysis

Sample:CE11123007-001  
Harvest/Lot ID: N/A  
Batch#: RDC-217031  
Metric Source Package #: N/A  
Batch Date: N/A  
Sample Size Received: 15 gram  
Total Weight/Volume: N/A  
Retail Product Size: N/A gram  
Ordered : 11/23/21  
sampled : 11/23/21  
Completed: 11/30/21 Expires: 11/30/22  
Sampling Method: SOP-024

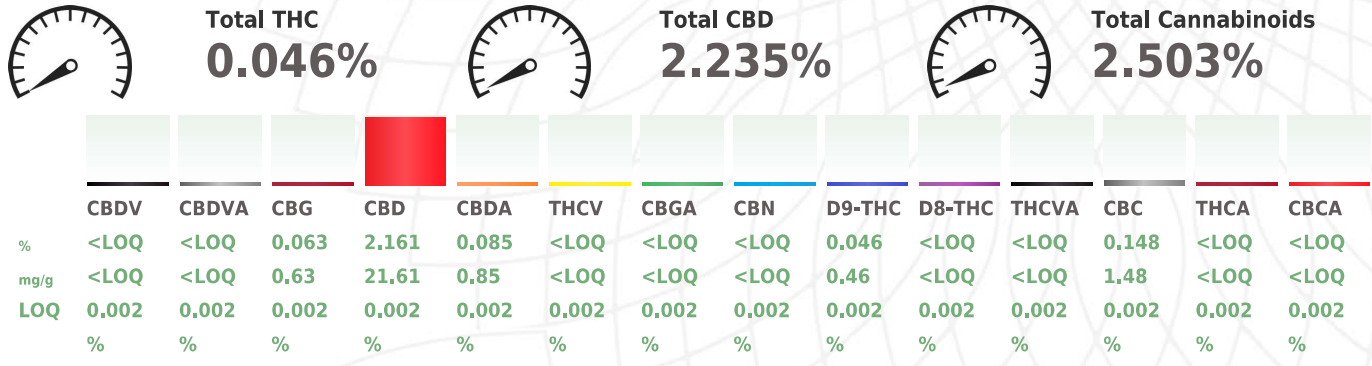
Nov 30, 2021 | Indomira/Green Earth Medicinals

License # R&D  
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 NOT TESTED	 Terpenes NOT TESTED

## CANNABINOID RESULTS



## Cannabinoid Profile Test

Analyzed by 11	Weight 0.939g	Extraction date : 11/29/21 03:11:28	Extracted By : 13
Analysis Method -SOP.T.40.020, SOP.T.30.050	Reviewed On - 11/30/21 16:34:10	Batch Date : 11/29/21 15:20:03	
Analytical Batch -CE000578POT	Instrument Used : HPLC 2030 EID 005 - Low Concentration	Running On :	

Reagent	Dilution	Consums. ID	Consums. ID
102721.R03	820	D01493069	F148560
111721.01		32009E-1232	0325891
020521.19		436020160AS3 436020338AS2 436021005AS3	
		C0000642	
		041CD-041C	
		042C4-042AL	

"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.

This report shall not be reproduced, unless in its entirety, without written approval from Kaycha Labs. This report is an 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. 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 #  
010-10166277B9D  
ISO Accreditation # 99861



Signature

11/30/21

Signed On



## POTENCY BATCH QC REPORT

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### 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	%

Analytical Batch - CE000578POT

Instrument Used : HPLC 2030 EID 005 - Low Concentration



### LCS

Cannabinoid	LOQ	Recovery	Units	Recovery Limits
CBG_WET	0.002	100.2	%	70-130
CBD_WET	0.002	98.9	%	70-130
CBDa_WET	0.002	100.5	%	70-130
THCV_WET	0.002	101.3	%	70-130
CBGA_WET	0.002	96.9	%	70-130
CBN_WET	0.002	99.6	%	70-130
D9-THC_WET	0.002	99.3	%	70-130
CBC_WET	0.002	99.5	%	70-130
THCA_WET	0.002	98	%	70-130
CBC-A_WET	0.002	107	%	70-130

Analytical Batch - CE000578POT

Instrument Used : HPLC 2030 EID 005 - Low Concentration



## Certificate of Analysis

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

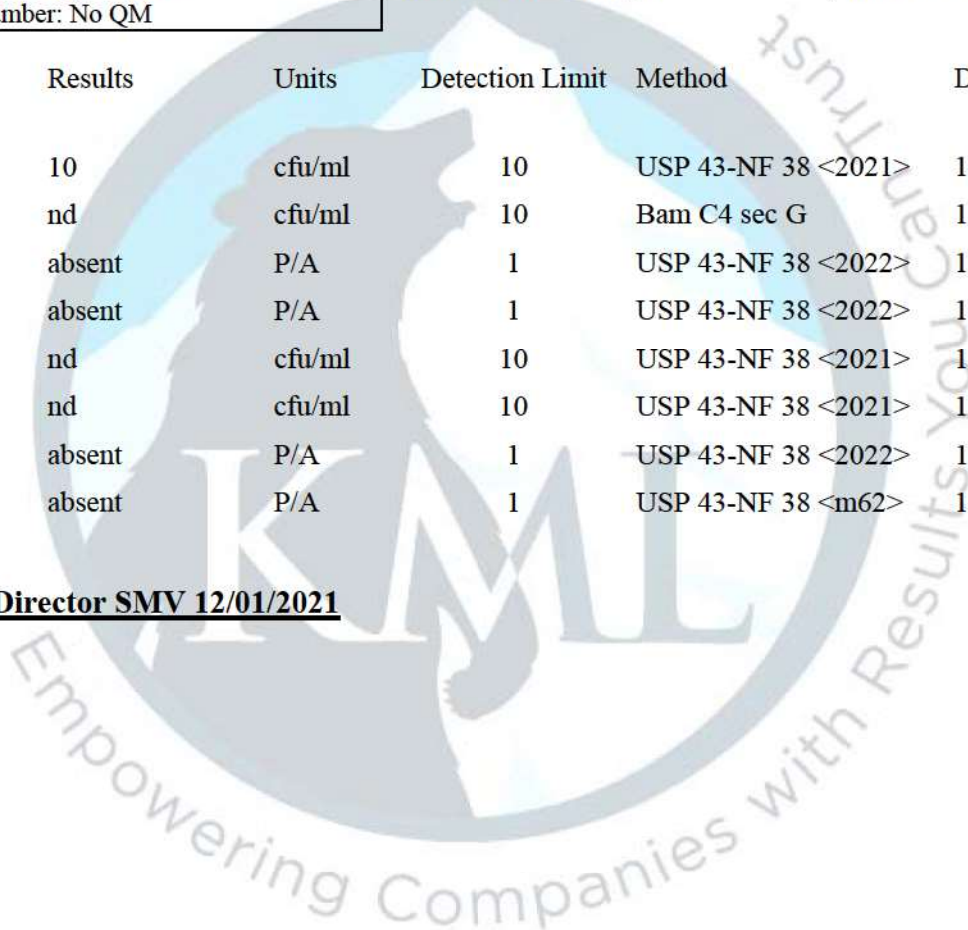
Invoice Number: 21.1483  
 PO Number: 11/23/21  
 Received Date: 11/26/2021  
 Number of Samples: 01  
 Project Name: Routine Testing

### Microbiology Report:

Lab #: 21-12395	Sample Lot: RDC-217031	Sample Date: 11/22/2021
Sample Name: CBD Oral Drops Regular	Additional ID: Strength	Plated Date: 11/26/2021
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>	11/29/2021
Coliforms	nd	cfu/ml	10	Bam C4 sec G	11/27/2021
<i>E. coli</i>	absent	P/A	1	USP 43-NF 38 <2022>	11/29/2021
<i>Staph aureus</i>	absent	P/A	1	USP 43-NF 38 <2022>	11/29/2021
Yeast	nd	cfu/ml	10	USP 43-NF 38 <2021>	12/01/2021
Mold	nd	cfu/ml	10	USP 43-NF 38 <2021>	12/01/2021
Salmonella	absent	P/A	1	USP 43-NF 38 <2022>	11/30/2021
<i>Pseudo. aeruginosa</i>	absent	P/A	1	USP 43-NF 38 <m62>	11/29/2021

**Approved By: QA Director SMV 12/01/2021**



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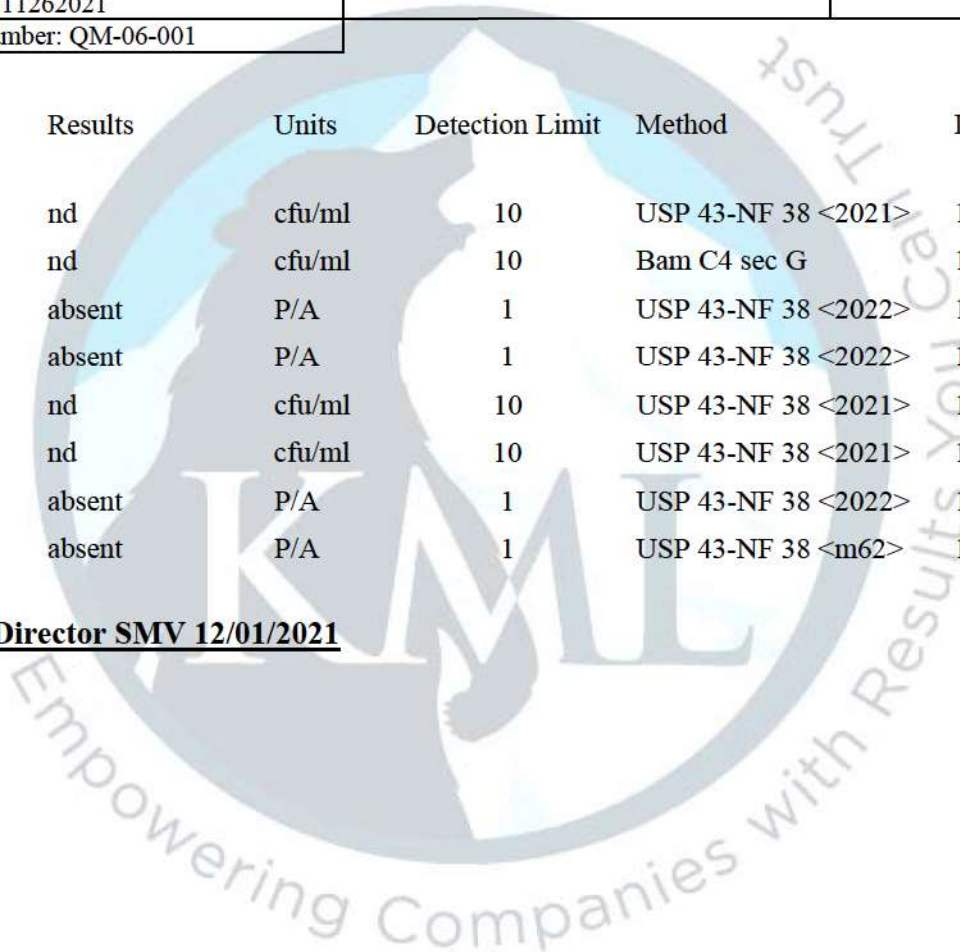
Invoice Number: 21.1483  
 PO Number: 11/23/21  
 Received Date: 11/26/2021  
 Number of Samples: 01  
 Project Name: Routine Testing

### Microbiology Report:

Lab #: Control 11262021	Additional ID: Negative Control Purposes	Plated Date: 11/26/2021
Sample Name: Control 11262021		
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>	11/29/2021
Coliforms	nd	cfu/ml	10	Bam C4 sec G	11/27/2021
<i>E. coli</i>	absent	P/A	1	USP 43-NF 38 <2022>	11/29/2021
<i>Staph aureus</i>	absent	P/A	1	USP 43-NF 38 <2022>	11/29/2021
Yeast	nd	cfu/ml	10	USP 43-NF 38 <2021>	12/01/2021
Mold	nd	cfu/ml	10	USP 43-NF 38 <2021>	12/01/2021
Salmonella	absent	P/A	1	USP 43-NF 38 <2022>	11/30/2021
<i>Pseudo. aeruginosa</i>	absent	P/A	1	USP 43-NF 38 <m62>	11/29/2021

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