

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

# Certificate of Analysis

### Kaycha Labs 画微器画

CBD Extra Oral Drops Spearmint N/A

Sample Type: Ingestible

Sample: CE20504001-001 Harvest/Lot ID: N/A Batch#: EDC-228022

Metrc Source Package #: N/A

Metrc #: N/A

Batch Date: N/A Sample Size Received: 10 gram

Total Weight/Volume: N/A Retail Product Size: N/A gram

ordered: 05/04/22 sampled: 05/04/22

Completed: 05/06/22 Sampling Method: SOP-024

Page 1 of 2

# May 06, 2022 | Indomira/Green Earth Medicinals

License # R&D 2305 Ashland St, Ste C360 Ashland, OR, 97520, US



SAFETY RESULTS











Microbials



Residuals Solvents



Filth



Water Activity







Terpenes NOT TESTED

**TESTED** 



Cannabinoid



0.1311%



5.1894%



**Total Cannabinoids** 5.8042%



540, 14, 11, 12

Analysis Method -SOP.T.40.020, SOP.T.30.050

Reviewed On - 05/05/22 12:58:41 Analytical Batch -CE001060POT

Batch Date: 05/04/22 12:28:04

0.951g

Instrument Used: HPLC 2030 EID 005 - Low Concentration

Running On: 05/04/22 14:17:50

Dilution: 820

Reagent: 032922.R01: 040822.04: 120920.02

Consumables: 21/07/20; 210407; 031022-A; ASC000G11324BSF; 12315-120CC-120D; 933C4-933AL; 00321166--6 00280879 00321305-4 00321165-6 00322250-6; 2132 81421

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

05/04/22 12:31:03

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** 

State License # 010-10166277B9D ISO Accreditation # 99861

05/06/22

Signed On

Signature



Central Point, OR, 97502, US

#### Kaycha Labs

CBD Extra Oral Drops Spearmint

Sample Type : Ingestible



## POTENCY BATCH QC REPORT

Page 2 of 2



#### **METHOD BLANK**

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

Analytical Batch - CE001060POT

Instrument Used: HPLC 2030 EID 005 - Low Concentration



#### **LCS**

Cannabinoid	LOQ	Recovery	Units	Recovery Limits
CBG_WET	0.002	105.3	%	80-120
CBD_WET	0.002	98.7	%	90-110
CBDA_WET	0.002	94.9	%	90-110
CBGA_WET	0.002	98.5	%	80-120
CBN_WET	0.002	98.1	%	80-120
D9-THC_WET	0.002	97.9	%	90-110
D8-THC_WET	0.002	105.5	%	90-110
CBC_WET	0.002	98.2	%	80-120
THCA_WET	0.002	96.4	%	90-110
CBC-A_WET	0.002	97.4	%	80-120

Analytical Batch - CE001060POT

Instrument Used: HPLC 2030 EID 005 - 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. 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=Nto Tebeteted, NA=Not Analyzed, ppm=Parts Per Million, pplb=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** 

State License # 010-10166277B9D ISO Accreditation # 99861

Signature

05/06/22

Signed On

# 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: 22.0622 PO Number: 05/03/22 Received Date: 05/05/2022 Number of Samples: 01

Project Name: Routine Testing

#### **Microbiology Report:**

Lab #: 22-4262	Sample Lot: EDC-228022	Sample Date: 05/03/2022
Sample Name: CBD Extra Sublingual Drops	Additional ID:	Plated Date: 05/05/2022
Qualifying Material Number: No OM		

Test Performed	Results	Units	Detection Limit	Method	Date Analyzed
Aerobic Plate Count	nd	cfu/ml	10	USP 43-NF 38 <2021>	05/08/2022
Coliforms	nd	cfu/ml	10	Bam C4 sec G	05/06/2022
E. coli	absent	P/A	1	USP 43-NF 38 <2022>	05/08/2022
Staph aureus	absent	P/A	1	USP 43-NF 38 <2022>	05/08/2022
Yeast	nd	cfu/ml	10	USP 43-NF 38 <2021>	05/10/2022
Mold	nd	cfu/ml	10	USP 43-NF 38 <2021>	05/10/2022
Salmonella	absent	P/A	1	USP 43-NF 38 <2022>	05/09/2022
Pseudo. aeruginosa	absent	P/A	1	USP 43-NF 38 <m62></m62>	05/08/2022

300 Mering Companies with Res Approved By: QA Director SMV 05/10/2022



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 ‡.

AT-1805

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: 22.0622 PO Number: 05/03/22 Received Date: 05/05/2022 Number of Samples: 01

Project Name: **Routine Testing** 

#### **Microbiology Report:**

Lab #: Control 05052022	Additional ID: Negative Control Purposes	Plated Date: 05/05/2022
Sample Name: Control 05052022		
Qualifying Matarial Number QM 06 001		

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>	05/08/2022
Coliforms	nd	cfu/ml	10	Bam C4 sec G	05/06/2022
E. coli	absent	P/A	1	USP 43-NF 38 <2022>	05/08/2022
Staph aureus	absent	P/A	1	USP 43-NF 38 <2022>	05/08/2022
Yeast	nd	cfu/ml	10	USP 43-NF 38 <2021>	05/10/2022
Mold	nd	cfu/ml	10	USP 43-NF 38 <2021>	05/10/2022
Salmonella	absent	P/A	1	USP 43-NF 38 <2022>	05/09/2022
Pseudo. aeruginosa	absent	P/A	1	USP 43-NF 38 <m62></m62>	05/08/2022

200 Mering Companies with Approved By: QA Director SMV 05/10/202



Malter

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