

Certificate of Analysis

Jan 28, 2022 | Planta Rx®

1205 71st St.

Miami Beach, FL 33141



Kaycha Labs

Olive Oil 1000MG

Matrix: Derivative



Sample: KN20118009-001 Harvest/Lot ID: 00I010122

> Batch#: 0010101220001 Seed to Sale# N/A Batch Date: 01/01/22

Sample Size Received: 240 ml Total Weight/Volume: N/A

Retail Product Size: 240 ml Ordered: 01/10/22

sampled: 01/10/22

Completed: 01/28/22 Expires: 01/28/23 Sampling Method: SOP Client Method

PASSED

PRODUCT IMAGE







PASSED





PASSED



PASSED



Mycotoxins

PASSED



Solvents

PASSED



PASSED



Water Activity





Moisture

NOT TESTED



NOT TESTED

CANNABINOID RESULTS



Total THC TOTAL THC/Bottle :0 mg



0.481%

Total Cannabinoids 0.481%Total Cannabinoids/Bottle :1062.048 mg



₩ Fi	lth		PASSEI
Analyzed By	Weight	Extraction date	Extracted By
1692	0.5247q	01/24/22	1692
Analyte	LOD	Pass/Fail	Result
Filth and Foreign M	laterial 0.3	Pass	ND
Analysis Method	-SOP.T.40.013	Batch Date: 01/24	/22 12:46:13
Analytical Batch	-KN001852FIL	Reviewed On - 01/	24/22 14:10:48
Instrument Used	: E-AMS-138 I	Microscope	
Running On :			
This includes but is n manufacturing waste	ot limited to hair, and by-products	insects, feces, packaging c . A SW-2T13 Stereo Microsc	ontaminants, and ope is use for inspection

Cannabinoid Profile Test

	racted By :
113 1.033g 01/18/22 03:01:29 113	
Analysis Method - Expanded Measurement of Uncertainty: Flower Matrix d9-THC:12.7%, THCa: 9.5%, TOTAL THC 11. 1%. These uncertainties Reviewed On - 01/19/22	
represent an expanded uncertainty expressed at approximately the 95% confidence level using a coverage factor k=2 for a normal distribution. 13:38:00 Batch Date: 01/18/22 1: Analytical Batch LYMON 31900T Instrument Island - 1801 C. E-SHLOND 8. Bunning On 1	10:29:32

947B9291.217

This report shall not be reproduced, unless in its entirety, without written approval from Kaycha Labs. This report is 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 for human safety for consumption and/or inhalation. The result >99% are 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 F.S. Rule 64-4.310.

Sue Ferguson

State License # n/a ISO Accreditation # 17025:2017



01/28/22

Signature



Kaycha Labs

Olive Oil 1000MG

Matrix : Derivative



Certificate of Analysis

PASSED

1205 71st St, Miami Beach, FL 33141 **Telephone:** (786)-572-3034 Email: sales@cbdmiamishop.com Harvest/Lot ID: 00I010122

Batch#: 0010101220001 Sampled: 01/10/22 Ordered: 01/10/22

Sample Size Received: 240 ml Total Weight/Volume: N/A Completed: 01/28/22 Expires: 01/28/23 Sample Method: SOP Client Method

Page 2 of 4

PASSED

Pass/Fail Result



Pest	ticid	es
------	-------	----

_					
Pesticides	LOD	Units	Action Level	Pass/Fail	Res
ABAMECTIN B1A	0.01	ppm	0.3	PASS	ND
ACEPHATE	0.01	ppm	3	PASS	ND
ACEQUINOCYL	0.01	ppm	2	PASS	ND
ACETAMIPRID	0.01	ppm	3	PASS	ND
ALDICARB	0.01	ppm	0.1	PASS	ND
AZOXYSTROBIN	0.01	ppm	3	PASS	ND
BIFENAZATE	0.01	ppm	3	PASS	ND
BIFENTHRIN	0.01	ppm	0.5	PASS	ND
BOSCALID	0.01	ppm	3	PASS	ND
CARBARYL	0.01	ppm	0.5	PASS	ND
CARBOFURAN	0.01	ppm	0.1	PASS	ND
CHLORANTRANILIPROLE	0.01	ppm	3	PASS	ND
CHLORMEQUAT CHLORIDE	0.01	ppm	3	PASS	ND
CHLORPYRIFOS	0.01	ppm	0.1	PASS	ND
CLOFENTEZINE	0.01	ppm	0.5	PASS	ND
COUMAPHOS	0.01	ppm	0.1	PASS	ND
CYPERMETHRIN	0.01	ppm	1	PASS	ND
DAMINOZIDE	0.01	ppm	0.1	PASS	ND
DIAZANON	0.01	ppm	0.2	PASS	ND
DICHLORVOS	0.01	ppm	0.1	PASS	ND
DIMETHOATE	0.01	ppm	0.1	PASS	ND
DIMETHOMORPH	0.01	ppm	3	PASS	ND
ETHOPROPHOS	0.01	ppm	0.1	PASS	ND
ETOFENPROX	0.01	ppm	0.1	PASS	ND
ETOXAZOLE	0.01	ppm	1.5	PASS	ND
FENHEXAMID	0.01	ppm	3	PASS	ND
FENOXYCARB	0.01	ppm	0.1	PASS	ND
FENPYROXIMATE	0.01	ppm	2	PASS	ND
FIPRONIL	0.01	ppm	0.1	PASS	ND
FLONICAMID	0.01	ppm	2	PASS	ND
FLUDIOXONIL	0.01	ppm	3	PASS	ND
HEXYTHIAZOX	0.01	ppm	2	PASS	ND
IMAZALIL	0.01	ppm	0.1	PASS	ND
IMIDACLOPRID	0.01	ppm	3	PASS	ND
KRESOXIM-METHYL	0.01	ppm	1	PASS	ND
MALATHION	0.01	ppm	2	PASS	ND
METALAXYL	0.01	ppm	3	PASS	ND
METHIOCARB	0.01	ppm	0.1	PASS	ND
METHOMYL	0.01	ppm	0.1	PASS	ND
MEVINPHOS	0.01	ppm	0.1	PASS	ND
MYCLOBUTANIL	0.01	ppm	3	PASS	ND
NALED	0.01	ppm	0.5	PASS	ND
OXAMYL	0.01	ppm	0.5	PASS	ND
PACLOBUTRAZOL	0.01	ppm	0.1	PASS	ND
PERMETHRINS	0.01	ppm	1	PASS	ND
PHOSMET	0.01	ppm	0.2	PASS	< 0.0
FIIOSPIEI	0.01	Phili	0.2		~0.0

0.01 PIPERONYL BUTOXIDE ppm PASS 0.01 0.4 ND PRALLETHRIN ppm PASS PROPICONAZOLE 0.01 ppm ND PASS PROPOXUR 0.01 ND mag 0.1 0.01 PASS ND PYRETHRINS mag PASS **PYRIDABEN** 0.01 ND ppm PASS SPINETORAM 0.01 mag ND 0.01 PASS ND SPIROMESIEEN ppm PASS SPIROTETRAMAT 0.01 ppm ND PASS SPIROXAMINE 0.01 ppm 0.1 ND **TEBUCONAZOLE** 0.01 mag PASS ND THIACI OPRID 0.01 ppm 0.1 PASS ND THIAMETHOXAM 0.01 ppm PASS ND

0.01 ppm

0.01

LOD Units

Level

0

TOTAL SPINOSAD

TRIFLOXYSTROBIN

Pesticides

PASS

PASS

Batch Date: 01/24/22 09:59:05

ND

PASSED

Extraction date Extracted By Analyzed by 143 0.5197g 01/24/22 1 Analysis Method - SOP.T.30.060, SOP.T.40.060, 01/24/22 11:01:43 Analytical Batch - KN001848PES Reviewed On 01/24/22 14:10:48 Instrument Used: E-SHI-125 Pesticides

Reagent 010722.R03 051021.01 011822.R09 011922.R16 011922.R15 010622.R02

Running On: 01/24/22 16:25:38

Dilution Consumables ID 200618634 947.271

Pesticide screen is performed using LC-MS which can screen down to below single digit ppb concentrations for regulated Pesticides. Currently we analyze for 57 Pesticides. (Method: SOP.T.30.060 Sample Preparation for Pesticides Analysis via LCMSMS and SOP.T40.060 Procedure for Pesticide Quantification Using LCMS). Analytes ISO pending. *Based on FL action limits. *

This report shall not be reproduced, unless in its entirety, without written approval from Kaycha Labs. This report is 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 for human safety for consumption and/or inhalation. The result >99% are 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 F.S. Rule 64-4.310.

Sue Ferguson

State License # n/a ISO Accreditation # 17025:2017



01/28/22

Signature



Kaycha Labs

Olive Oil 1000MG

Matrix : Derivative



Certificate of Analysis

PASSED

Planta Rx®

1205 71st St, Miami Beach, FL 33141 Telephone: (786)-572-3034 Email: sales@cbdmiamishop.com Sample : KN20118009-001 Harvest/Lot ID: 00I010122

Batch#:0010101220001 Sampled:01/10/22 Ordered:01/10/22 Sample Size Received: 240 ml Total Weight/Volume: N/A Completed: 01/28/22 Expires: 01/28/23 Sample Method: SOP Client Method

Page 3 of 4



Residual Solvents

PASSED

Solvent	LOD	Units	Action Level	Pass/Fail	Result
PROPANE	500	ppm	2100	PASS	ND
BUTANES (N-BUTANE)	500	ppm	2000	PASS	ND
METHANOL	25	ppm	3000	PASS	ND
ETHYLENE OXIDE	0.5	ppm	5	PASS	ND
PENTANES (N-PENTANE)	75	ppm	5000	PASS	ND
ETHANOL	500	ppm	5000	PASS	ND
ETHYL ETHER	50	ppm	5000	PASS	ND
1.1-DICHLOROETHENE	0.8	ppm	8	PASS	ND
ACETONE	75	ppm	5000	PASS	ND
2-PROPANOL	50 6	ppm	500	PASS	ND
ACETONITRILE	6	ppm	410	PASS	ND
DICHLOROMETHANE	12.5	ppm	600	PASS	ND
N-HEXANE	25	ppm	290	PASS	ND
ETHYL ACETATE	40	ppm	5000	PASS	ND
CHLOROFORM	0.2	ppm	60	PASS	ND
BENZENE	0.1	ppm	2	PASS	ND
1,2-DICHLOROETHANE	0.2	ppm	5	PASS	ND
HEPTANE	500	ppm	5000	PASS	ND
TRICHLOROETHYLENE	2.5	ppm	80	PASS	ND
TOLUENE	15	ppm	890	PASS	ND
TOTAL XYLENES - M, P & O - DIMETHYLBENZENE	15	ppm	2170	PASS	ND



Residual Solvents

PASSED

Analyzed by

Weight 0.02847g

Extraction date 01/24/22 03:01:58

Extracted By

Analysis Method -SOP.T.40.032 Analytical Batch -KN001851SOL

Instrument Used: E-SHI-106 Residual Solvents

Running On: 01/24/22 16:34:17 Batch Date: 01/24/22 11:29:17 Reviewed On - 01/25/22 18:14:59

Reagent

Dilution

Consumables ID

R2017.062 G201-062

Residual solvents screening is performed using GC-MS which can detect below single digit ppm concentrations. Currently we analyze for 22 residual solvents. (Method: SOP.T.40.032 Residual Solvents Analysis via GC-MS). Analytes ISO pending. *Based on FL action limits.

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 for human safety for consumption and/or inhalation. The result >99% are 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 F.S. Rule 64-4.310.

Sue Ferguson

Lab Directo

State License # n/a ISO Accreditation # 17025:2017 Sulinguan

01/28/22

Signature



Kaycha Labs

Olive Oil 1000MG

N/A

Matrix : Derivative



Certificate of Analysis

PASSED

Planta Rx®

1205 71st St, Miami Beach, FL 33141 Telephone: (786)-572-3034 Email: sales@cbdmiamishop.com Sample : KN20118009-001 Harvest/Lot ID: OOI010122

Batch#: 0010101220001 Sampled: 01/10/22 Ordered: 01/10/22 Sample Size Received: 240 ml Total Weight/Volume: N/A Completed: 01/28/22 Expires: 01/28/23 Sample Method: SOP Client Method

Page 4 of 4



Microbials

PASSED



Mycotoxins

PASSED

Analyte	LOD	Result	Pass / Fail
LISTERIA MONOCYTOGENE		not present in 1 gram.	PASS
ESCHERICHIA COLI SHIGELLA SPP		not present in 1 gram.	PASS
SALMONELLA SPECIFIC GENE		not present in 1 gram.	PASS
ASPERGILLUS FLAVUS		not present in 1 gram.	PASS
ASPERGILLUS FUMIGATUS		not present in 1 gram.	PASS
ASPERGILLUS NIGER		not present in 1 gram.	PASS
ASPERGILLUS TERREUS		not present in 1 gram.	PASS

Analysis Method -SOP.T.40.043

Analytical Batch -KN001846MIC Batch Date: 01/24/22 08:02:03

Instrument Used: Micro E-HEW-069

Running On:

Analyzed by	Weight	Extraction date	Extracted By
1692	1.0129g	NA	NA
Reagent			Dilution

030121.01 122921.01 121521.04 030421.09

Microbiological testing for Fungal and Bacterial Identification via Polymerase Chain Reaction (PCR) method consisting of sample DNA amplified via tandem Polymerase Chain Reaction (PCR) as a crude lysate which avoids purification. (Method SOP.T.40.043) If a pathogenic Escherichia Coli, Salmonella, Aspergillus fumigatus, Aspergillus flavus, Aspergillus niger, or Aspergillus terreus is detected in 1g of a sample, the sample fails the microbiological-impurity testing.

Analyte	LOD	Units	Result	Pass / Fail	Action Level
AFLATOXIN G2	0.002	ppm	ND	PASS	0.02
AFLATOXIN G1	0.002	ppm	ND	PASS	0.02
AFLATOXIN B2	0.002	ppm	ND	PASS	0.02
AFLATOXIN B1	0.002	ppm	ND	PASS	0.02
OCHRATOXIN A+	0.002	ppm	ND	PASS	0.02
TOTAL MYCOTOXINS	0.002	ppm	ND	PASS	

Analysis Method -SOP.T.30.060, SOP.T.40.060

Analytical Batch -KN001849MYC | Reviewed On - 01/25/22 10:48:01

Instrument Used: E-SHI-125 Mycotoxins

Running On: 01/24/22 16:29:24 | Batch Date: 01/24/22 09:59:49

Analyzed by	Weight	Extraction date	Extracted By
143	0.5197g	01/24/22 04:01:28	143

Aflatoxins B1, B2, G1, G2, and Ochratoxins A testing using LC-MS. (Method: SOP.T.30.060 for Sample Preparation and SOP.T40.060 Procedure for Mycotoxins Quantification Using LCMS. LOQ 1.0 ppb). Total Aflatoxins (Aflotoxin B1, B2, G1, G2) must be $<\!20\mu g/Kg$. Ochratoxins must be $<\!20\mu g/Kg$. Analytes ISO pending. *Based on FL action limits.



Heavy Metals

PASSED

Metal	LOD	Unit	Result	Pass / Fail	Action Level	
ARSENIC-AS	0.02	ppm	ND	PASS	1.5	
CADMIUM-CD	0.02	ppm	ND	PASS	0.5	
MERCURY-HG	0.02	ppm	ND	PASS	3	
LEAD-PB	0.02	ppm	ND	PASS	0.5	

Analyzed by Weight Extraction date Extracted By 12 225g NA NA

Analysis Method -SOP.T.40.050, SOP.T.30.052

Analytical Batch -KN001853HEA | Reviewed On - 01/25/22 18:16:20

Instrument Used : Metals ICP/MS

Running On: | Batch Date: 01/24/22 13:44:40

Dilution	Consums. ID
1	7226/0030021
	12235-110CD-110C
	Dilution

Heavy Metals screening is performed using ICP-MS (Inductively Coupled Plasma – Mass Spectrometer) which can screen down to below single digit ppb concentrations for regulated heavy metals using Method SOP.T.30.052 Sample Preparation for Heavy Metals Analysis via ICP-MS and SOP.T.40.050 Heavy Metals Analysis via ICP-MS.

Analysis via ICP-MS. Analysis via ICP-MS. 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, NN=Not Analyzed, ppm=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 for human safety for consumption and/or inhalation. The result > 99% are 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 F.S. Rule 64-4.310.

Sue Ferguson

Lab Director

State License # n/a ISO Accreditation # 17025:2017



01/28/22

Signature