

Certificate of Analysis

Apr 21, 2022 | Green Roads

5150 SW 48TH WAY Davie, FL, 33314, US



Kaycha Labs

SWEET SLEEP CBD+CBN OIL

Matrix: Edible



Sample: KN20419011-002 Harvest/Lot ID: D14Y02

Batch#: BMR0112/GRW0103 Seed to Sale# N/A

Batch Date: 04/14/22

Sample Size Received: 34.8 gram Total Weight/Volume: N/A

Retail Product Size: 34.8 gram

ordered: 04/18/22 sampled: 04/18/22

Completed: 04/21/22 Sampling Method: SOP Client Method

PASSED

Page $1 ext{ of } 4$

SAFETY RESULTS PRODUCT IMAGE











PASSED





Residuals **PASSED** Solvents PASSED



PASSED









PASSED



Cannabinoid

CBN 0.2225%

CBN/Bottle: 77.43 mg



Total CBD

Total CBD/Bottle: 742.214 mg



Total Cannabinoids

(ii)

Total Cannabinoids/Bottle: 819.644 mg



W F	iitn		PASSEL
Analyzed By	Weight	Extraction date	Extracted By
1692	0.522g	04/19/22	1692
Analyte	LO	D Pass/Fail	Result
Filth and Foreign N	laterial 0.3	Pass	ND
Analysis Method	-SOP.T.40.0	13 Batch Date: 04/19	9/22 09:08:21
Analytical Batch	-KN002283F	L Reviewed On - 04	19/22 12:36:41
Instrument Used	: E-AMS-138	Microscope	
Running On:			
Running On :			

Cannabinoid Profile Test

atch Date: 04/19/22 10:24:32

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 Direc

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



Signature

04/21/22



Kaycha Labs

SWEET SLEEP CBD+CBN OIL

Matrix : Edible



Certificate of Analysis

PASSED

5150 SW 48TH WAY Davie, FL, 33314, US Telephone: (844) 747-3367

Email: LAURA@GREENROADSWORLD.COM

Harvest/Lot ID: D14Y02

Sampled: 04/18/22 Odered: 04/18/22

Batch#: BMR0112/GRW0103 Sample Size Received: 34.8 gram Total Weight/Volume: N/A **Completed:** 04/21/22 **Expires:** 04/21/23 Sample Method: SOP Client Method

Page 2 of 4



Pesticides

5	PASSED

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
ENPYROXIMATE	0.01	ppm	2	PASS	ND
FIPRONIL	0.01	mag	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
	0.01		3	PASS	ND
METALAXYL	0.01	ppm	0.1	PASS	ND
METHIOCARB		ppm		PASS	
METHOMYL	0.01	ppm	0.1		ND ND
MEVINPHOS	0.01	ppm	0.1	PASS	
		ppm	3	PASS	ND
				PASS	ND
NALED	0.01	ppm	0.5		
MYCLOBUTANIL NALED OXAMYL	0.01	ppm	0.5	PASS	ND
NALED OXAMYL PACLOBUTRAZOL	0.01 0.01 0.01	ppm ppm	0.5 0.1	PASS PASS	ND
NALED OXAMYL	0.01	ppm	0.5	PASS	

Pesticides	LOD	Units	Action Level	Pass/Fail	Result	
PIPERONYL BUTOXIDE	0.01	ppm	3	PASS	ND	
PRALLETHRIN	0.01	ppm	0.4	PASS	ND	
PROPICONAZOLE	0.01	ppm	1	PASS	ND	
PROPOXUR	0.01	ppm	0.1	PASS	ND	
PYRETHRINS	0.01	ppm	1	PASS	ND	
PYRIDABEN	0.01	ppm	3	PASS	ND	
SPINETORAM	0.01	ppm	3	PASS	ND	
SPIROMESIFEN	0.01	ppm	3	PASS	ND	
SPIROTETRAMAT	0.01	ppm	3	PASS	ND	
SPIROXAMINE	0.01	ppm	0.1	PASS	ND	
TEBUCONAZOLE	0.01	ppm	1	PASS	ND	
THIACLOPRID	0.01	ppm	0.1	PASS	ND	
THIAMETHOXAM	0.01	ppm	1	PASS	ND	
TOTAL SPINOSAD	0.01	ppm	3	PASS	ND	
TRIFLOXYSTROBIN	0.01	ppm	3	PASS	ND	

Pesticides

PASSED

Extracted by:

Analysis Method -SOP.T.30.060, SOP.T.40.060 Analytical Batch -KN002291PES

Instrument Used : E-SHI-125 Pesticides Running on:

Weight: **Extraction date:**

Reviewed On: 04/21/22 16:30:51 Batch Date: 04/20/22 10:23:51

Analyzed by: Dilution: 1

Pesticide analysis is performed using LC-MSMS which can quantify down to below single digit ppb concentrations for regulated Pesticides. Currently we analyze for 61 Pesticides. (Methods: SOP.T.30.065 Sample Preparation for Pesticides Analysis via LCMSMS and SOP.T40.065 Procedure for Pesticide Quantification Using LCMSMS). *Based on FL action limits. *

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

Instrument Used : Running on : Analyzed by:

Weight:

Extraction date:

Reviewed On: Batch Date:

Extracted by:

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



Signature

04/21/22



Kaycha Labs

SWEET SLEEP CBD+CBN OIL

Matrix : Edible



Certificate of Analysis

PASSED

5150 SW 48TH WAY Davie, FL, 33314, US Telephone: (844) 747-3367 Email: LAURA@GREENROADSWORLD.COM Harvest/Lot ID: D14Y02

Batch#: BMR0112/GRW0103 Sample Size Received: 34.8 gram Sampled: 04/18/22 Odered: 04/18/22

Total Weight/Volume: N/A **Completed:** 04/21/22 **Expires:** 04/21/23 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	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 138

Weight

0.02983g

Extraction date 04/20/22 12:04:08 Extracted By

Analysis Method -SOP,T.40.032 Analytical Batch - KN002280SOL

Instrument Used: E-SHI-106 Residual Solvents

Running On:

Batch Date: 04/19/22 08:37:29

Reviewed On - 04/21/22 19:25:21

Dilution: 1 Reagent:

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

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

04/21/22

Signature



Kaycha Labs

SWEET SLEEP CBD+CBN OIL

Matrix : Edible



Certificate of Analysis

PASSED

5150 SW 48TH WAY Davie, FL, 33314, US Telephone: (844) 747-3367

Email: LAURA@GREENROADSWORLD.COM

Harvest/Lot ID: D14Y02

Sampled: 04/18/22

Odered: 04/18/22

Batch#: BMR0112/GRW0103 Sample Size Received: 34.8 gram Total Weight/Volume: N/A

Completed: 04/21/22 Expires: 04/21/23 Sample Method: SOP Client Method

Page 4 of 4



Microbials



Mycotoxins

PASSED

Analyte	LOD	Result	Pass / Fail	Action Level
LISTERIA MONOCYTOGENE	2000	ND	PASS	2000
ESCHERICHIA COLI SHIGELLA SPP	1726	ND	PASS	1726
SALMONELLA SPECIFIC GENE	10000	ND	PASS	10000
ASPERGILLUS FLAVUS	10000	ND	PASS	10000
ASPERGILLUS FUMIGATUS	10000	ND	PASS	10000
ASPERGILLUS NIGER	10000	ND	PASS	10000
ASPERGILLUS TERREUS	10000	ND	PASS	10000
TOTAL YEAST AND MOLD	10	<10	PASS	100000

Analysis Method - SOP.T.40.043 Analytical Batch - KN002281MIC Instrument Used: Micro E-HEW-069

Running on

Analyzed by: Weight: 1692 1.0083q

Batch Date: 04/19/22 08:47:28 Extraction date: 04/19/22 12:04:59

Reviewed On: 04/21/22 19:25:06

Reviewed On: 04/20/22 17:12:10

Batch Date: 04/19/22 09:06:13

Extracted by: 1692

Dilution: 1

Reagent: 030121.01: 121721.06: 122021.01

Consumables:

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

Analysis Method - SOP.T.40.043 Analytical Batch - KN002282TYM Instrument Used: E-HEW-069
Running on:

Analyzed by: 1692 **Weight:** 1.0083g Extraction date: 04/19/22 12:04:16

Dilution: 1 Reagent: 030121.01 Consumables:

Total yeast and mold testing is performed utilizing MPN and traditional culture based techniques in accordance with F.S. Rule 64ER20-39.

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

					Fall	Level	
AFLATOXIN G	2	0.00)2 ppn	n ND	PASS	0.02	
AFLATOXIN G	1	0.00)2 ppn	n ND	PASS	0.02	
AFLATOXIN B	2	0.00)2 ppn	n ND	PASS	0.02	
AFLATOXIN B	1	0.00)2 ppn	n ND	PASS	0.02	
OCHRATOXIN	A+	0.00	2 ppn	n ND	PASS	0.02	
TOTAL MYCOT	TOXINS	0.00)2 ppn	n ND	PASS	0.02	
		G 2 - G					

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

Analytical Batch -KN002292MYC | Reviewed On - 04/21/22 19:25:44

Instrument Used:

Running On: | Batch Date: 04/20/22 10:24:21

Weight **Extraction date Extracted By** Analyzed by 20g NA

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 <20µg/Kg. Ochratoxins must be <20µg/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 da	ate	Extr	acted B	у

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

Analytical Batch -KN002284HEA | Reviewed On - 04/20/22 19:38:27

Instrument Used: Metals ICP/MS

Running On: | Batch Date: 04/19/22 10:24:12

Dilution: 1 Reagent:

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.

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. This report shall not be reproduced, unless in its entirety, without written approval from Kaycha Labs. This report is

Sue Ferguson

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

Signature

04/21/22