

Certificate of Analysis

Jan 18, 2022 | Bad Days

Rochester, NY, 14624, US



Kaycha Labs

BD Sour Apple Off Hours

Matrix: Derivative



Sample: KN20111003-004 Harvest/Lot ID: BD-OH-SA-RnD

> Batch#: 1-3-22 Seed to Sale# N/A

Batch Date: 01/03/22

Sample Size Received: 60 gram

Total Weight/Volume: N/A Retail Product Size: 6.2 gram

Ordered: 01/06/22

sampled: 01/06/22 Completed: 01/18/22 Expires: 01/18/23

Sampling Method: SOP Client Method

PASSED

Page 1 of 4

PRODUCT IMAGE

SAFETY RESULTS





Heavy Metals **PASSED**



Microbials **PASSED**



Residuals **PASSED** Solvents



PASSED PASSED



Water Activity

(iii)



NOT TESTED



MISC.

NOT TESTED

CANNABINOID RESULTS



Total THC



Total CBD TOTAL CBD/Chewable :5.27 mg



Total Cannabinoids

Total Cannabinoids/Chewable :21.018 mg



	-IIth			PASSE
		Extraction	Extracted	
Analyzed By	Weight	date	Ву	
1692	0.5312g	01/13/22		1692
Analyte		LOD	A.L	Result
Filth and Foreign	Material	0.3	3	ND
Analysis Metho	od -SOP.T.40.013	Batch Date	e: 01/13/22 1	12:51:27
Analytical Bato	h -KN001805FIL	Reviewed	On - 01/13/22	2 14:43:04
Instrument Use	ed: E-AMS-138 M	icroscope		
Running On:				
This includes but is a	ont limited to hair insect	r force parkaning	contaminants an	d manufacturing was

Cannabinoid Profile Test

Analysis Method -E Reviewed On -01/12/22 08:37:29 Batch Date: 01/11/22 09:35:07

Consums, ID 081321.R04

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

Lab Director

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



01/18/22

Signature



Kaycha Labs

BD Sour Apple Off Hours





Certificate of Analysis

PASSED

350 Buell Road

Rochester, NY, 14624, US Telephone: (315) 406-6767 Email: seth@nowave.com

Sample: KN20111003-004 Harvest/Lot ID: BD-OH-SA-RnD

Batch#: 1-3-22 Sampled: 01/06/22 Ordered: 01/06/22

Sample Size Received: 60 gram Total Weight/Volume: N/A

Completed: 01/18/22 Expires: 01/18/23 Sample Method: SOP Client Method

Page 2 of 4



Pesticides

PASSED

Pesticides	LOD	Units	Action Level	Resu
BAMECTIN B1A	0.01	ppm	0.3	ND
СЕРНАТЕ	0.01	ppm	3	ND
CEQUINOCYL	0.01	ppm	2	ND
CETAMIPRID	0.01	ppm	3	ND
LDICARB	0.01	ppm	0.1	ND
ZOXYSTROBIN	0.01	ppm	3	ND
FENAZATE	0.01	ppm	3	ND
FENTHRIN	0.01	ppm	0.5	ND
OSCALID	0.01	ppm	3	ND
ARBARYL	0.01	ppm	0.5	ND
ARBOFURAN	0.01	ppm	0.1	ND
HLORANTRANILIPROLE	0.01	ppm	3	ND
HLORMEQUAT CHLORIDE	0.01	ppm	3	ND
HLORPYRIFOS	0.01	ppm	0.1	ND
OFENTEZINE	0.01	ppm	0.5	ND
DUMAPHOS	0.01	ppm	0.1	ND
PERMETHRIN	0.01	ppm	1	ND
AMINOZIDE	0.01	ppm	0.1	ND
AZANON	0.01	ppm	0.2	ND
CHLORVOS	0.01	ppm	0.1	ND
METHOATE	0.01	ppm	0.1	ND
METHOMORPH	0.01	ppm	3	ND
THOPROPHOS	0.01	ppm	0.1	ND
TOFENPROX	0.01	ppm	0.1	ND
TOXAZOLE	0.01	ppm	1.5	ND
ENHEXAMID	0.01	ppm	3	ND
ENOXYCARB	0.01	ppm	0.1	ND
ENPYROXIMATE	0.01	ppm	2	ND
PRONIL	0.01	ppm	0.1	ND
ONICAMID	0.01	ppm	2	ND
UDIOXONIL	0.01	ppm	3	ND
EXYTHIAZOX	0.01	ppm	2	ND
IAZALIL	0.01	ppm	0.1	ND
IIDACLOPRID	0.01	ppm	3	ND
RESOXIM-METHYL	0.01	ppm	1	ND
ALATHION	0.01	ppm	2	ND
ETALAXYL	0.01	ppm	3	ND
ETHIOCARB	0.01	ppm	0.1	ND
ETHOMYL	0.01	ppm	0.1	ND
EVINPHOS	0.01	ppm	0.1	ND
YCLOBUTANIL	0.01	ppm	3	0.436
ALED	0.01	ppm	0.5	ND
XAMYL	0.01	ppm	0.5	ND
ACLOBUTRAZOL	0.01	ppm	0.1	ND
ERMETHRINS	0.01	ppm	1	0.085
HOSMET	0.01	ppm	0.2	0.063 ND

Pesticides	LOD	Units	Action Level	Result
PIPERONYL BUTOXIDE	0.01	ppm	3	< 0.05
PRALLETHRIN	0.01	ppm	0.4	ND
PROPICONAZOLE	0.01	ppm	1	ND
PROPOXUR	0.01	ppm	0.1	ND
PYRETHRINS	0.01	ppm	1	ND
PYRIDABEN	0.01	ppm	3	ND
SPINETORAM	0.01	ppm	3	ND
SPIROMESIFEN	0.01	ppm	3	ND
SPIROTETRAMAT	0.01	ppm	3	ND
SPIROXAMINE	0.01	ppm	0.1	ND
TEBUCONAZOLE	0.01	ppm	1	ND
THIACLOPRID	0.01	ppm	0.1	ND
THIAMETHOXAM	0.01	ppm	1	ND
TOTAL SPINOSAD	0.01	ppm	3	ND
TRIFLOXYSTROBIN	0.01	ppm	3	ND

Analyzed by	Weight	Extraction date	Extracted By
143	0.5548g	01/13/22 03:01:06	143
Analysis Method - SOP.	T.30.060, SOP.T.40.060	.// // /	
Analytical Batch - KN001801PES			Reviewed On- 01/13/22 14:43:04
Instrument Used : E-SH			
Running On: 01/13/22	12:10:15		Batch Date: 01/13/22 08:29:29
Reagent		Dilution	Consums. ID
010722.R03		10	200618634
051021.03			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.T.40.060 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 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 Director

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

01/18/22

Signature



Kaycha Labs

BD Sour Apple Off Hours

Matrix : Derivative



Certificate of Analysis

PASSED

350 Buell Road

Rochester, NY, 14624, US Telephone: (315) 406-6767 Email: seth@nowave.com

Sample: KN20111003-004 Harvest/Lot ID: BD-OH-SA-RnD

Batch#: 1-3-22 Sampled: 01/06/22 Ordered: 01/06/22

Sample Size Received: 60 gram Total Weight/Volume: N/A

Completed: 01/18/22 Expires: 01/18/23 Sample Method: SOP Client Method

Page 3 of 4



Residual Solvents

PASSED



Residual Solvents



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 & - DIMETHYLBENZENE	0 15	ppm	2170	PASS	ND

		7/9	L	412.
L	Analyzed by	Weight	Extraction date	Extracted B

Analysis Method -SOP.T.40.032

Analytical Batch -KN001813SOL Reviewed On - 01/18/22 17:53:53

Instrument Used: E-SHI-106 Residual Solvents

Running On: 01/14/22 16:38:00 Batch Date: 01/14/22 09:22:11

Reagent Dilution Consums. ID

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 Director

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



01/18/22

Signature



Kaycha Labs

BD Sour Apple Off Hours

Matrix : Derivative



Certificate of Analysis

PASSED

350 Buell Road

Rochester, NY, 14624, US Telephone: (315) 406-6767 Email: seth@nowave.com

Sample: KN20111003-004 Harvest/Lot ID: BD-OH-SA-RnD

Batch#: 1-3-22 Sampled: 01/06/22 Ordered: 01/06/22

Result

not present in 1 gram. not present in 1 gram. not present in 1 gram. not present in 1 gram.

not present in 1 gram.

not present in 1 gram.

not present in 1 gram.

Sample Size Received: 60 gram Total Weight/Volume: N/A

Completed: 01/18/22 Expires: 01/18/23 Sample Method: SOP Client Method

Page 4 of 4



Microbials

PASSED



OCHRATOXIN A+

TOTAL MYCOTOXINS

Mycotoxins

PASSED

Analyte
LISTERIA MONOCYTOGENE
ESCHERICHIA COLI SHIGELLA SPP
SALMONELLA SPECIFIC GENE
ASPERGILLUS FLAVUS
ASPERGILLUS FUMIGATUS
ASPERGILLUS NIGER
ACDED CILLUC TERREILS

Analysis Method -SOP.T.40.043

Analytical Batch -KN001808MIC Batch Date: 01/13/22 15:51:23

Instrument Used: Micro E-HEW-069

Running On: Analyzed by

Allulyzcu	IJ y	
1692		

Weight 1.0205g

Extraction da

LOD

Analyte	LOD	Units	Result	Action Level
AFLATOXIN G2	0.002	ppm	ND	0.02
AFLATOXIN G1	0.002	ppm	ND	0.02
AFLATOXIN B2	0.002	ppm	ND	0.02
AFLATOXIN B1	0.002	ppm	ND	0.02

ppm

mag

ND

ND

0.02

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

Analytical Batch -KN001802MYC | Reviewed On - 01/14/22 13:57:07

0.002

0.002

Instrument Used: E-SHI-125 Mycotoxins Running On: 01/13/22 12:10:23

Batch Date: 01/13/22 08:30:09

ite	Extracted By		
	NA		14

nalyzed by Weight **Extraction date Extracted By** 0.5548a 01/13/22 04:01:32 143

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 <20µg/Kg. Ochratoxins must be <20µg/Kg. Analytes ISO pending. *Based on FL action limits.

Dilution

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 figer, or Aspergillus trreus is detected in 1g of a sample, the sample fails the microbiological-impurity testing.



Heavy Metals

PASSED

Reagent	Dilution	Consums. ID
121421.05	50	7226/0030021
011022.R08		210221060
080421.R13		
011022.R07		

Metal	LOD	Unit	Result	Action Level
ARSENIC-AS	0.02	ppm	ND	1.5
CADMIUM-CD	0.02	ppm	ND	0.5
MERCURY-HG	0.02	ppm	ND	3
LEAD-PB	0.02	ppm	ND	0.5
Analyzed by	Weight	Extraction	date	Extracted By
12	0.2543g	01/14/22 02:	01:19	12

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

Analytical Batch -KN001807HEA | Reviewed On - 01/14/22 15:13:33

Instrument Used : Metals ICP/MS

Running On:

Batch Date: 01/13/22 13:54:06

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.

Sue Ferguson

Lab Director

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



01/18/22

Signature