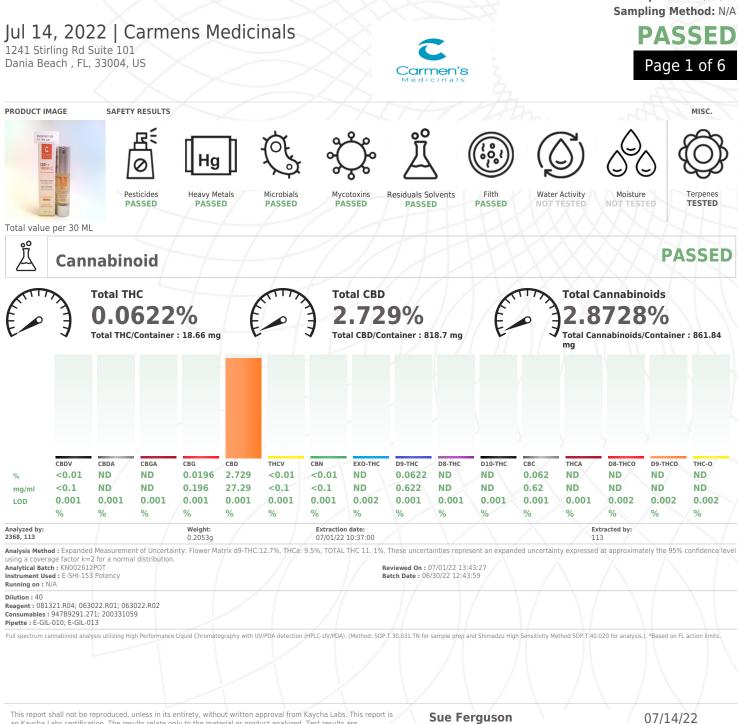


Certificate of Analysis

Sample:KN20630007-004 Harvest/Lot ID: F13EA Batch#: F13EA Seed to Sale# N/A Batch Date: 06/27/22 Sample Size Received: 30 ml Total Batch Size: N/A Retail Product Size: 30 ml Ordered : 06/27/22 Sampled : 06/27/22 Completed: 07/14/22



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

Signature



Kaycha Labs 🔳 🏾 🛣 🕍 🔳

750 Mg CBD Serum N/A Matrix : Derivative



PASSED

Certificate of Analysis

Carmens Medicinals

1241 Stirling Rd Suite 101 Dania Beach , FL, 33004, US Telephone: (888) 328-6445 Email: info@carmensmedicinals.com Sample : KN20630007-004 Harvest/Lot ID: F13EA Batch# : F13EA Sampled : 06/27/22 Ordered : 06/27/22

Sample Size Received : 30 ml Total Batch Size : N/A Completed : 07/14/22 Expires: 07/14/23 Sample Method : SOP Client Method



TESTED

٩

Terpenes

Terpenes	LOD (%)	mg/ml	%	Result (%)	Terpenes		LOD (%)	mg/ml	%	Result (%)
RANS-CARYOPHYLLENE	0.007	ND	ND		HEXAHYDROTHYMOL		0.007	ND	ND	
GUAIOL	0.007	<0.2	< 0.02		EUCALYPTOL		0.007	ND	ND	
IMONENE	0.007	3.99	0.399		ISOBORNEOL		0.007	ND	ND	
INALOOL	0.007	ND	ND		FARNESENE		0.007	ND	ND	
IEROL	0.007	ND	ND		FENCHONE		0.007	ND	ND	
CIMENE	0.007	ND	ND		GAMMA-TERPINENE		0.007	ND	ND	
ALPHA-PHELLANDRENE	0.007	ND	ND		GERANIOL		0.007	ND	ND	
ULEGONE	0.007	ND	ND		Analyzed by:	Weight:	E	traction d	ate:	Extracted by:
ABINENE	0.007	ND	ND		2368, 138, 12	1.0249g		7/01/22 11		138
SABINENE HYDRATE	0.007	ND	ND		Analysis Method : SOP.T.40	0.090				
ERPINEOL	0.007	ND	ND		Analytical Batch : KN00261					On: 07/13/22 11:40:58
ERPINOLENE	0.007	ND	ND		Instrument Used : E-SHI-10 Running on : N/A	9 Terpenes		E	Batch Date	a: 07/01/22 10:45:56
GERANYL ACETATE	0.007	ND	ND		Dilution : 10					
RANS-NEROLIDOL	0.007	ND	ND		Reagent : 092221.02					
ALENCENE	0.007	ND	ND		Consumables : 294108110	; n/a; 211214634-D	; 947B92	91.271		
SOPULEGOL	0.007	ND	ND		Pipette : N/A					V X N X I
ALPHA-HUMULENE	0.007	ND	ND		Terpenoid profile screening is 38 terpenes using Method SOI	performed using GC-I	45 with Liq	uid Injection	n (Gas Chro	matography – Mass Spectrometer) which can scre
LPHA-PINENE	0.007	ND	ND		So terpenes using method sol	.1.40.050 Terpendid	Analysis vi	a ocimis. Ai	larytes 150	rending
LPHA-TERPINENE	0.007	ND	ND							
BETA-MYRCENE	0.007	ND	ND							
ETA-PINENE	0.007	ND	ND							
BORNEOL	0.013	ND	ND							
AMPHENE	0.007	ND	ND							
AMPHOR	0.013	ND	ND							
ARYOPHYLLENE OXIDE	0.007	ND	ND							
EDROL	0.007	ND	ND							
ALPHA-BISABOLOL	0.007	0.373	0.0373							
LPHA-CEDRENE	0.007	ND	ND							
CIS-NEROLIDOL	0.007	ND	ND							
	0.007	ND	ND							
3-CARENE										
3-CARENE FENCHYL ALCOHOL	0.007	ND	ND							

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, pm=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 Lucinguson Signature 07/14/22



Kaycha Labs 🔳 🛞 🗠 🖉

..... 750 Mg CBD Serum N/A Matrix : Derivative



PASSED

Certificate of Analysis

Carmens Medicinals

1241 Stirling Rd Suite 101 Dania Beach , FL, 33004, US Telephone: (888) 328-6445 Email: info@carmensmedicinals.com

Sample : KN20630007-004 Harvest/Lot ID: F13EA Batch# : F13EA Sampled : 06/27/22 Ordered : 06/27/22

Sample Size Received : 30 ml Total Batch Size : N/A Completed : 07/14/22 Expires: 07/14/23 Sample Method : SOP Client Method

Page 3 of 6

PASSED

£: 0

Pesticides

Pesticide	LOD	Units	Action Level	Pass/Fail	Result
ABAMECTIN B1A	0.01	ppm	0.3	PASS	ND
АСЕРНАТЕ	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
ZOXYSTROBIN	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	mag	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
TOFENPROX	0.01	mag	0.1	PASS	ND
TOXAZOLE	0.01	ppm	1.5	PASS	ND
ENHEXAMID	0.01	ppm	3	PASS	ND
ENOXYCARB	0.01	ppm	0.1	PASS	ND
	0.01	ppm	2	PASS	ND
FIPRONIL	0.01	ppm	0.1	PASS	ND
FLONICAMID	0.01	ppm	2	PASS	ND
FLUDIOXONIL	0.01		3	PASS	ND
	0.01	ppm	2	PASS	ND
HEXYTHIAZOX	0.01	ppm	2	PASS	ND
MAZALIL		ppm	3	PASS	
MIDACLOPRID	0.01	ppm	-		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
DXAMYL	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	ND

Pesticide	28	LOD	Units	Action Level	Pass/Fail	Result
PIPERONYL BUTO	XIDE	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
TRIFLOXYSTROBI	N	0.01	ppm	3	PASS	ND
Analyzed by: 12	Weight: 0.2035g	Extraction 07/14/22 17			Extracted 12	by:
Analytical Batch :	:SOP.T.30.060, SOP.T. KN002659PES :E-SHI-125 Pesticides	40.060		d On :07/14/ ate :07/14/22		
Reagent : N/A						

Consumables : N/A

Pipette : N/A

Protect : N/A 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.

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 Detectod, NA=Not Analyzed, pm=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 Dire

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

Sulinguar Signature

07/14/22



Kaycha Labs

750 Mg CBD Serum N/A Matrix : Derivative



PASSED

Certificate of Analysis

Carmens Medicinals

1241 Stirling Rd Suite 101 Dania Beach , FL, 33004, US Telephone: (888) 328-6445 Email: info@carmensmedicinals.com Sample : KN20630007-004 Harvest/Lot ID: F13EA Batch# : F13EA Sampled : 06/27/22 Ordered : 06/27/22

Sample Size Received : 30 ml Total Batch Size : N/A Completed : 07/14/22 Expires: 07/14/23 Sample Method : SOP Client Method



PASSED

Page 4 of 6

Residual Solvents

olvents	LOD	Units	Action Level	Pass/Fail	Result
ROPANE	500	ppm	2100	PASS	ND
UTANES (N-BUTANE)	500	ppm	2000	PASS	ND
ETHANOL	25	ppm	3000	PASS	ND
THYLENE OXIDE	0.5	ppm	5	PASS	ND
ENTANES (N-PENTANE)	75	ppm	5000	PASS	ND
THANOL	500	ppm	5000	PASS	ND
THYL ETHER	50	ppm	5000	PASS	ND
1-DICHLOROETHENE	0.8	ppm	8	PASS	ND
CETONE	75	ppm	5000	PASS	ND
PROPANOL	50	ppm	500	PASS	ND
CETONITRILE	6	ppm	410	PASS	ND
ICHLOROMETHANE	12.5	ppm	600	PASS	ND
HEXANE	25	ppm	290	PASS	ND
THYL ACETATE	40	ppm	5000	PASS	ND
HLOROFORM	0.2	ppm	60	PASS	ND
ENZENE	0.1	ppm	2	PASS	ND
2-DICHLOROETHANE	0.2	ppm	5	PASS	ND
EPTANE	500	ppm	5000	PASS	ND
RICHLOROETHYLENE	2.5	ppm	80	PASS	ND
DLUENE	15	ppm	890	PASS	ND
OTAL XYLENES - M, P & O - DIMETHYLBENZENE	15	ppm	2170	PASS	ND
nalyzed by: Wei A N/A	ight:	Extraction date: N/A		Extracted by: N/A	
Analysis Method : SOP.T.40.032 Analytical Batch : KN002606SOL nstrument Used : E-SHI-106 Residual Solvents Aunning on : N/A			Reviewed On : 07/01/22 Batch Date : 06/29/22 1		X X

Consumables : R2017.126; G201.126

Pipette : N/A

Residual solvents analysis 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). *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, pm=Parts Per Million, ppb=Parts Per Billion. Limit to 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 Signature

07/14/22



Kaycha Labs 🔳 🛞 🗠 🛓

750 Mg CBD Serum N/A Matrix : Derivative



PASSED

Certificate of Analysis

Carmens Medicinals

1241 Stirling Rd Suite 101 Dania Beach , FL, 33004, US **Telephone:** (888) 328-6445 **Email:** info@carmensmedicinals.com Sample : KN20630007-004 Harvest/Lot ID: F13EA Batch# : F13EA Sampled : 06/27/22 Ordered : 06/27/22

Sample Size Received : 30 ml Total Batch Size : N/A Completed : 07/14/22 Expires: 07/14/23 Sample Method : SOP Client Method

Mvcotoxins

ഹ്ം

Page	5	of	6	

PASSED

Analyte		LOD	Units	Result	Pass / Fail	Acti Lev
LISTERIA MO	NOCYTOGENE			Not Present	PASS	
ESCHERICHIA SPP	COLI SHIGELLA			Not Present	PASS	
SALMONELLA	SPECIFIC GENE			Not Present	PASS	
ASPERGILLUS	FLAVUS			Not Present	PASS	
ASPERGILLUS	FUMIGATUS			Not Present	PASS	
ASPERGILLUS	NIGER			Not Present	PASS	
ASPERGILLUS	TERREUS			Not Present	PASS	
Analyzed by: 2368, 1692, 12	Weight: 1.0374g		tion date: 22 08:47:52		Extracted 1692	by:
	d: SOP.T.40.043 1: KN002615MIC d: Micro E-HEW-069			Dn : 07/06/22 : 07/01/22 08		

Pipette : N/A

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.

مکو		~~~~~				- 3	8
Analyte			LOD	Units	Result	Pass / Fail	Action Level
AFLATOXIN O	52		0.002	ppm	ND	PASS	0.02
AFLATOXIN O	G1 00		0.002	ppm	ND	PASS	0.02
AFLATOXIN E	32		0.002	ppm	ND	PASS	0.02
AFLATOXIN E	31		0.002	ppm	ND	PASS	0.02
OCHRATOXIN	IA+		0.002	ppm	ND	PASS	0.02
TOTAL MYCO	TOXINS		0.002	ppm	ND	PASS	0.02
Analyzed by: 12	Weight: 22g	Extract N/A	tion date		Extr N/A	acted by:	
Analytical Batc	d:SOP.T.30.060, SOP h:KN002655MYC ed:E-SHI-125 Mycoto /A				:07/14/22)7/14/22 1		H
Dilution : N/A Reagent : N/A Consumables : Pipette : N/A	N/A						

Aflatoxins B1, B2, G1, G2, and Ochratoxins A testing using LC-MS. (Method: SOP.T.30.060 for Sample Preparation and SOP.T40.065 Procedure for Mycotoxins Quantification Using LCMSMS. LOQ 5.0 ppb). *Based on FL action limits.

[нд] Н	eavy M	letals		X	PAS	SED
Metal	71-1	LOD	Units	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: 2368, 138, 12	Weight: 0.2714g	Extraction date 07/05/22 14:04			Extracted	by:
Analysis Method : SC Analytical Batch : KN Instrument Used : Me Running on : N/A	002623HEA	Reviewe		06/22 15: 5/22 09:42		
Dilution : 50 Reagent : N/A Consumables : N/A Pipette : N/A	X				X	7

Heavy Metals screening is performed using ICP-MS (Inductively Coupled Plasma – Mass Spectrometer) which can screen down to single digit ppb concentrations for regulated heavy metals using Method SOP.T.30.082 Sample Preparation for Heavy Metals Analysis via ICP-MS and SOP.T.40.082TN 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 Detectod, NA=Not Analyzed, pm=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

Sulinguse Signature

07/14/22



Kaycha Labs 📷 🛞 🗠 🖄

750 Mg CBD Serum N/A Matrix : Derivative



PASSED

Page 6 of 6

Certificate of Analysis

Carmens Medicinals

1241 Stirling Rd Suite 101 Dania Beach , FL, 33004, US **Telephone:** (888) 328-6445 **Email:** info@carmensmedicinals.com Sample : KN20630007-004 Harvest/Lot ID: F13EA Batch# : F13EA Sampled : 06/27/22 Ordered : 06/27/22

PASSED

Sample Size Received : 30 ml Total Batch Size : N/A Completed : 07/14/22 Expires: 07/14/23 Sample Method : SOP Client Method



Filth/Foreign Material

Analyte Filth and Foreign	Material	LOD 1		Result ND	P/F PASS	Action Leve
Analyzed by: 2368, 1692	Weight: 0.5097g		raction date: 30/22 16:44:07	7	Extr 1693	acted by: 2
Analysis Method : S Analytical Batch : K Instrument Used : E Running on : N/A	N002613FIL		Revie		06/30/22 16 5/30/22 13:3	
Dilution : N/A Reagent : N/A Consumables : N/A Pipette : N/A						

This includes but is not limited to hair, insects, feces, packaging contaminants, and manufacturing waste and by-products. A SW-2T13 Stereo Microscope is use for inspection.

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

huluquon Signature 07/14/22