

Certificate of Analysis

May 17, 2020 | PharmaCanna

2615 state Road 7 Wellington FL, United States 33414



Kaycha Labs

CBDream 300 Regular

Matrix: Edible



Sample: DA00511004-001 Harvest/Lot ID: B20D300R

> Seed to Sale #N/A Batch Date : N/A

Batch#: B20D300R Sample Size Received: 300 mg

> Retail Product Size: 0.421 Ordered: 05/07/20

Sampled: 05/07/20

Completed: 05/17/20 Expires: 05/17/21 Sampling Method: SOP Client Method

PASSED

Page 1 of 4

PRODUCT IMAGE

SAFETY RESULTS







PASSED



Heavy Metals PASSED



Microbials



Mycotoxins

Reviewed On - 05/15/20 17:53:45

Solvents **PASSED**



PASSED



Water Activity



Moisture **NOT TESTED**



MISC.

NOT TESTED

CANNABINOID RESULTS



Total THC 0.000%THC/Capsule :0.000 mg

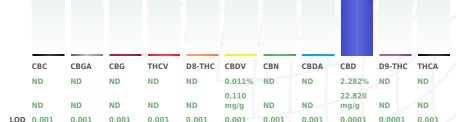


Total CBD 2.282% CBD/Capsule:9.607 mg



Total Cannabinoids

Total Cannabinoids/Capsule :9.654 mg





Filth

PASSED

Weight Extraction date Analyzed By LOD(ppm) Extracted By 1q NA

Analysis Method -SOP.T.40.013 Batch Date: 05/15/20 10:22:53 Analytical Batch -DA012463FIL Reviewed On - 05/15/20 10:23:41 Instrument Used: Filth/Foreign Material Microscope

Cannabinoid Profile Test

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

Analyzed by Weight Extraction date: Extracted By:

Analytical Batch - DA012393POT Instrument Used: DA-LC-003 Batch Date: 05/13/20 11:51:01 Dilution 280678841 914C4-914AK 929C6-929H 032320.27 051120.R07

051120.R06 Full spectrum cannabinoid analysis utilizing High Performance Liquid Chromatography with UV detection (HPLC-UV). (Method: SOP.T.30.050 for sample prep and Shimadzu High Sensitivity Method SOP.T.40.020 for analysis. LOQ for all cannabinoids is 1 mg/L).

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.

Jorge Segredo Lab Director

State License # n/a ISO Accreditation # 97164



05/17/2020

Signed On Signature



Kaycha Labs

CBDream 300 Regular

Matrix: Edible



Certificate of Analysis

PASSED

2615 state Road 7 Wellington FL, United States 33414 Telephone: 9543050078 **Email:** johnny@pharmacanna.us Sample: DA00511004-001 Harvest/LOT ID: B20D300R

Batch#: B20D300R Sampled: 05/07/20 Ordered: 05/07/20

Sample Size Received: 300 mg Completed: 05/17/20 Expires: 05/17/21 Sample Method: SOP Client Method

Page 2 of 4



Pesticides

PASSED

Pesticides	LOD	Units	Action Level	Result
ABAMECTIN B1A	0.01	ppm	0.3	ND
ACEPHATE	0.01	ppm	3	ND
ACEQUINOCYL	0.01	ppm	2	ND
ACETAMIPRID	0.01	ppm	3	ND
ALDICARB	0.01	ppm	0.1	ND
AZOXYSTROBIN	0.01	ppm	3	ND
BIFENAZATE	0.01	ppm	3	ND
BIFENTHRIN	0.01	ppm	0.5	ND
BOSCALID	0.01	PPM	3	ND
CARBARYL	0.05	ppm	0.5	ND
CARBOFURAN	0.01	ppm	0.1	ND
CHLORANTRANILIPROLE	0.1	ppm	3	ND
CHLORMEQUAT CHLORIDE	0.05	ppm	3	ND
CHLORPYRIFOS	0.01	ppm	0.1	ND
CLOFENTEZINE	0.02	ppm	0.5	ND
COUMAPHOS	0.01	ppm	0.1	ND
DAMINOZIDE	0.01	ppm	0.1	ND
DIAZANON	0.01	ppm	0.2	ND
DICHLORVOS	0.01	ppm	0.1	ND
CYPERMETHRIN	0.05	ppm	1	ND
DIMETHOATE	0.01	ppm	0.1	ND
DIMETHOMORPH	0.02	ppm	3	ND
ETHOPROPHOS	0.01	ppm	0.1	ND
ETOFENPROX	0.01	ppm	0.1	ND
ETOXAZOLE	0.01	ppm	1.5	ND
FENHEXAMID	0.01	ppm	3	ND
FENOXYCARB	0.01	ppm	0.1	ND
FENPYROXIMATE	0.01	ppm	2	ND
FIPRONIL	0.01	ppm	0.1	ND
FLONICAMID	0.01	ppm	2	ND
FLUDIOXONIL	0.01	ppm	3	ND
HEXYTHIAZOX	0.01	ppm	2	ND
IMAZALIL	0.01	ppm	0.1	ND
IMIDACLOPRID	0.04	ppm	3	ND
KRESOXIM-METHYL	0.01	ppm	1	ND
MALATHION	0.02	ppm	2	ND
METALAXYL	0.01	ppm	3	ND
METHIOCARB	0.01	ppm	0.1	ND
METHOMYL	0.01	ppm	0.1	ND
METHYL PARATHION	0.005	ppm	0.1	ND
MEVINPHOS	0.01	ppm	0.1	ND

Pesticides	LOD	Units	Action Level	Result
MYCLOBUTANIL	0.01	ppm	3	ND
NALED	0.025	ppm	0.5	ND
OXAMYL	0.05	ppm	0.5	ND
PACLOBUTRAZOL	0.01	ppm	0.1	ND
PHOSMET	0.01	ppm	0.2	ND
PIPERONYL BUTOXIDE	0.1	ppm	3	ND
PRALLETHRIN	0.01	ppm	0.4	ND
PROPICONAZOLE	0.01	ppm	1	ND
PROPOXUR	0.01	ppm	0.1	ND
PYRETHRIN I	0.01	ppm	1	ND
PYRETHRIN II	0.01	ppm	1	ND
PYRETHRINS	0.05	ppm	1	ND
PYRIDABEN	0.02	ppm	3	ND
SPINETORAM	0.02	PPM	3	ND
SPINOSAD (SPINOSYN A)	0.01	ppm	3	ND
SPINOSAD (SPINOSYN D)	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	0.05	ppm	1	ND
TOTAL CONTAMINANT LOAD (PESTICIDES)	0	PPM	20	ND
TOTAL PERMETHRIN	0.01	ppm	1	ND
TOTAL SPINOSAD	0.01	ppm	3	ND
TRIFLOXYSTROBIN	0.01	ppm	3	ND

PASSED **Pesticides** Ö

Extraction date Analyzed by Weight **Extracted By** 1.0498g 05/11/20 11:05:03

Analysis Method - SOP.T.30.065, SOP.T.40.065 , SOP.T.30.065, SOP.T40.070

Analytical Batch - DA012188PES Instrument Used : DA-LCMS-001_DER (PES) Batch Date : 05/05/20 12:48:04

Reviewed On- 05/15/20 10:23:41

Reagent Dilution Consums. ID

Pesticide screen is performed using LC-MS which can screen down to below single digit ppb concentrations for regulated Pesticides. Currently we analyze for 67 Pesticides. (Method: SOP.T.30.060 Sample Preparation for Pesticides Analysis via LCMSMS and SOP.T40.065 Procedure for Pesticide Quantification Using LCMS), * Volatile Pesticide Screening is performed using GC-MS which can screen down to below single digit ppb concentrations for regulated Pesticides. Analytes marked with an asterisk were tested using GC-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.

Jorge Segredo Lab Director

State License # n/a ISO Accreditation # 97164



05/17/2020

Signature Signed On



Kaycha Labs

CBDream 300 Regular

Matrix: Edible



Certificate of Analysis

PASSED

2615 state Road 7 Wellington FL, United States 33414 Telephone: 9543050078 Email: johnny@pharmacanna.us Sample: DA00511004-001 Harvest/LOT ID: B20D300R

Batch#: B20D300R Sampled: 05/07/20 Ordered: 05/07/20

Sample Size Received: 300 mg Completed: 05/17/20 Expires: 05/17/21 Sample Method: SOP Client Method

Page 3 of 4



Residual Solvents

PASSED



Analyzed by

Reagent

850

Residual Solvents



Solvent	LOD	Units	Action Level (PPM)	Pass/Fail	Result
1,1-DICHLOROETHENE	0.8	ppm	8	PASS	ND
1,2-DICHLOROETHANE	0.2	ppm	5	PASS	ND
2-PROPANOL	50	ppm	500	PASS	ND
ACETONE	75	ppm	5000	PASS	ND
ACETONITRILE	6	ppm	410	PASS	ND
BENZENE	0.1	ppm	2	PASS	ND
BUTANES (N-BUTANE)	500	ppm	2000	PASS	ND
CHLOROFORM	0.2	ppm	60	PASS	ND
DICHLOROMETHANE	12.5	ppm	600	PASS	ND
ETHANOL	500	ppm	5000	PASS	ND
ETHYL ACETATE	40	ppm	5000	PASS	ND
ETHYL ETHER	50	ppm	5000	PASS	ND
ETHYLENE OXIDE	0.5	ppm	5	PASS	ND
HEPTANE	500	ppm	5000	PASS	ND
METHANOL	25	ppm	3000	PASS	ND
N-HEXANE	25	ppm	290	PASS	ND
PENTANES (N-PENTANE)	75	ppm	5000	PASS	ND
PROPANE	500	ppm	2100	PASS	ND
TOLUENE	15	ppm	890	PASS	ND
TRICHLOROETHYLENE	2.5	ppm	80	PASS	ND
XYLENES-M&P (1,3&1,4- DIMETHYLBENZENE)	27	ppm	2170	PASS	ND
XYLENES-O (1,2- DIMETHYLBENZENE)	13.5	ppm	2170	PASS	ND

Weight **Extraction date** 05/12/20 04:05:39

Extracted By

Analysis Method -SOP.T.40.032 Analytical Batch -DA012369SOL

Instrument Used: DA-GCMS-002 Batch Date: 05/12/20 15:53:03

0.02320

Reviewed On - 05/14/20 10:21:51

850

Dilution Consums, ID 00279984 161291-1 24154107

Residual solvents screening is performed using GC-MS which can detect below single digit ppm concentrations. Currently we analyze for 21 Residual solvents. (Method: SOP.T.30.032 Residual Solvents Analysis via GC-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

Jorge Segredo Lab Director

State License # n/a ISO Accreditation # 97164



05/17/2020

Signature

Signed On



Kaycha Labs

CBDream 300 Regular

Matrix: Edible



Certificate of Analysis

PASSED

2615 state Road 7 Wellington FL, United States 33414 Telephone: 9543050078 Email: johnny@pharmacanna.us

Sample : DA00511004-001 Harvest/LOT ID: B20D300R

PASSED

Batch#: B20D300R Sampled: 05/07/20 Ordered: 05/07/20

Sample Size Received: 300 mg Completed: 05/17/20 Expires: 05/17/21 Sample Method: SOP Client Method

Page 4 of 4

ॐ

0 0				
Analyte	LOD	Units	Result	Action Level (PPM)
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
OCHRATOXIN A+	0.002	ppm	ND	0.02

Analysis Method -SOP.T.30.065, SOP.T.40.065

Analytical Batch -DA012189 | Reviewed On - 05/16/20 01:28:08

Mycotoxins

Instrument Used: DA-LCMS-001_DER (MYC)

Batch Date: 05/05/20 12:49:16

Analyzed by	Weight	Extr
585	1g	05/13

raction date **Extracted By** 1/20 06:05:32

Aflatoxins B1, B2, G1, G2, and Ochratoxins A testing using LC-MS. (Method: SOP.T.30.065 for Sample Preparation and SOP.T40.065 Procedure for Mycotoxins Quantification Using LCMS. LOQ 1.0 Rea ppb). Aflatoxin B1, B2, G1, and G2 must individually be <20ug/Kg. Ochratoxins must be <20ug/Kg.



ASPERGILLUS_FUMIGATUS

ASPERGILLUS_NIGER

ASPERGILLUS_TERREUS

Analyte ASPERGILLUS FLAVUS

Microbials

PASSED

		Res

not present in 1 gram. not present in 1 gram

Analytical Batch -DA012321MIC | Reviewed On - 05/13/20 18:39:38 Instrument Used: PathogenDX PCR_Array Scanner DA-111,PathogenDX PCR_DA-171

Batch Date: 05/11/20 10:22:25

Analysis Method -SOP.T.40.043 / SOP.T.40.045

ESCHERICHIA COLI SHIGELLA SPP

SALMONELLA_SPECIFIC_GENE

Analyzed by Weight Extraction date **Extracted By** 05/11/20 10:05:37

Reagent 022520.07 101519.12 Dilution

181019-274

Consums, ID

Reagent	Consums. ID
032720.220	918C4-918J
032720.146	914C4-914AK
022120.26	929C6-929H
022120.268	50AX26219
032720.92	19323
022120.185	23819111
022120.51	190611634
022120.212	
050530.07	

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.



0309 0505 0505

0507

Heavy Metals

PASSED

Dilution

agent	Reagent	
820.R01 920.01	042720.R36 101819.07	
520.R05		
520.R04		
720.R13		
F20 D02		

Metal	LOD	Unit	Result	Action Level (PPM
ARSENIC	0.02	PPM	ND	1.5
CADMIUM	0.02	PPM	ND	0.5
LEAD	0.05	PPM	ND	0.5
MERCURY	0.02	PPM	ND	3
Analyzed by	Weight	Extractio	n date	Extracted By
53	0.2743g	05/11/20 01	L:05:02	1022

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

Analytical Batch -DA012300HEA | Reviewed On - 05/12/20 16:20:25

Instrument Used : DA-ICPMS-001 Batch Date: 05/11/20 08:22:14

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 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 (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.

Jorge Segredo Lab Director

State License # n/a ISO Accreditation # 97164



05/17/2020

Signature

Signed On