Certificate: 1222

SMITHERS

CERTIFICATE OF ANALYSIS

License #: 00000020LCVT89602592

Hemp THCa Flower

Batch #: #16
Strain: #16 After Dark
Parent Batch #:
Sample Collected: 10/26/2023 09:15:00
Published: 11/01/2023

Sample ID: 2310SMAZ0223.0680
Amount Received: 5.2 g
Sample Type: Flower - Cured
Received: 10/31/2023

COMPLIANCE FOR RETAIL

Regulated Analytes

Cannabinoid Profile (Q3) **Tested**	Microbial Contaminants Not Tested	Residual Solvents Not Tested
Pesticides, Fungicides, and Growth Regulators Not Tested	Mycotoxins Not Tested	Heavy Metals Not Tested

Additional Analytes (Not Regulated)

Terpenes Total (Q3) Not Tested	Moisture Analysis (Q3) Not Tested	Water Activity (Q3) Not Tested
Filth & Foreign (Q3) Not Tested	Homogeneity (Q3) Not Tested	

12.989%	Total THC
0.047%	Total CBD
ND	CBN
ND	CBG
15.705%	Total Cannabinoids (Q3)

Ahmed Munshi

Technical Laboratory Director

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ilac-MRA

PJLA Testing
Accreditation #: 103104

Certificate: 1222

CERTIFICATE OF ANALYSIS

License #: 00000020LCVT89602592

Cannabinoid Profile

HPLC **Tested**

Sample Prep

Batch Date: 10/30/2023
SOP: 418.AZ
Batch Number: 262

Sample Analysis

Date: 11/01/2023
SOP: 417.AZ - HPLC
Sample Weight: 0.103 g
Volume: 40 mL

Analyte	LOD (mg/g)	LOQ (mg/g)	Dil.	Actual % (w/w)	mg/g	Qualifier
CBC	0.125	0.379	1	ND	ND	
CBD	0.125	0.379	1	ND	ND	
CBDA	0.125	0.379	1	0.054	0.536	
CBDV	0.125	0.379	1	ND	ND	
CBG	0.125	0.379	1	ND	ND	
CBGA	0.125	0.379	1	0.478	4.780	
CBN	0.125	0.379	1	ND	ND	
d8-THC	0.125	0.379	1	ND	ND	
d9-THC	0.125	0.379	1	0.271	2.712	
THCA	0.125	0.379	1	14.502	145.018	
THCV	0.125	0.379	1	ND	ND	

Cannabinoid Totals	Actual % (w/w)	mg/g	Qualifier
Total THC	12.989	129.893	
Total CBD	0.047	0.470	
Total Cannabinoids	15.705	157.046	Q3

Total THC = THC + (0.877 x THCA) and Total CBD = CBD + (0.877 x CBDA)
ND = Not Detected, NT = Not Tested, <LOQ = Below Limit of Quantitation

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Qualifier Legend

B1 *The target analyte detected in the calibration is at or above the limit of quantitation, but the sample result for potency testing, is below the limit of quantitation.*

B2 *The target analyte detected in the calibration blank, or the method blank is at or above the limit of quantitation, but the sample result when testing for pesticides, fungicides, herbicides, growth regulators, heavy metals, or residual solvents, is below the maximum allowable concentration for the analyte.*

D1 *The limit of quantitation and the sample results were adjusted to reflect sample dilution.*

I1 *The relative intensity of a characteristic ion in a sample analyte exceeded the acceptance with respect to the reference spectra, indicating interference.*

L1 *When testing for pesticides, fungicides, herbicides, growth regulators, heavy metals, or residual solvents, the percent recovery of a laboratory control sample is greater than the acceptance limits, but the sample's target analytes were not detected above the maximum allowable concentrations for the analytes in the sample.*

M1 *The recovery from the matrix spike was high, but the recovery from the laboratory control sample was within acceptance criteria.*

M2 *The recovery from the matrix spike was low, but the recovery from the laboratory control sample was within acceptance criteria.*

M3 *The recovery from the matrix spike was unusable because the analyte concentration was disproportionate to the spike level, but the recovery from the laboratory control sample was within acceptance criteria.*

M4 *The analysis of a spiked sample required a dilution such that the spike recovery calculation does not provide useful information, but the recovery from the associated laboratory control sample was within acceptance criteria.*

M5 *The analyte concentration was determined by the method of standard addition, in which the standard is added directly to the aliquots of the analyzed sample.*

M6 *A description of the variance is described in the final report of testing according to R9-17- 404.06(B)(3)(d)(ii).*

Q1 *Sample integrity was not maintained.*

Q2 *The sample is heterogeneous, and sample homogeneity could not be readily achieved using routine laboratory practices.*

Q3 *Testing result is for informational purposes only and cannot be used to satisfy dispensary testing requirements in R9-17-317.01(A) or labeling requirements in R9-17-317.*

R1 *The relative percent difference for the laboratory control sample and duplicate exceeded the limit, but the recovery was within acceptance criteria.*

R2 *The relative percent difference for a sample and duplicate exceeded the limit.*

V1 *The recovery from continuing calibration verification standards exceeded the acceptance limits, but the sample's target analytes were not detected above the maximum allowable for the analytes in the sample.*

Notes:

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