



### **CERTIFICATE OF ANALYSIS**

License #: 00000020LCVT89602592

# **Hemp THCa Flower**

Batch #: 21 Strain: 43 Slurrycane

Parent Batch #:

**Sample Collected:** 11/08/2023 08:31:00

Published: 11/13/2023

**Sample ID:** 2311SMAZ0285.0917

Amount Received: 2.8 g Sample Type: Flower - Cured

Received: 11/09/2023



21.765% Total THC

0.061%

Total CBD

ND

ND CBG

25.943%

# **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** 

Total Cannabinoids (Q3)

Ahmed Munshi

**Technical Laboratory Director** 

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#### Certificate: 1650

# **Cannabinoid Profile**

**HPLC** 

**Tested** 

## **Sample Prep**

**Batch Date:** 11/08/2023 **SOP:** 418.AZ **Batch Number:** 325

## **Sample Analysis**

Date: 11/09/2023 SOP: 417.AZ - HPLC Sample Weight: 0.107 g Volume: 40 mL

Analyte	LOD (mg/g)	LOQ (mg/g)	Dil.	Actual % (w/w)	mg/g	Qualifier
CBC	0.120	0.365	1	ND	ND	
CBD	0.120	0.365	1	ND	ND	
CBDA	0.120	0.365	1	0.070	0.699	
CBDV	0.120	0.365	1	ND	ND	
CBG	0.120	0.365	1	ND	ND	
CBGA	0.120	0.365	1	1.269	12.692	
CBN	0.120	0.365	1	ND	ND	
d8-THC	0.120	0.365	1	ND	ND	
d9-THC	0.120	0.365	1	0.153	1.527	
THCA	0.120	0.365	1	23.077	230.767	
THCV	0.120	0.365	1	ND	ND	

Cannabinoid Totals	Actual % (w/w)	mg/g	Qualifier	
Total THC	21.765	217.653		
Total CBD	0.061	0.613		
Total Cannabinoids	25.943	259.429	Q3	

Total THC = THC +  $(0.877 \times THCA)$  and Total CBD = CBD +  $(0.877 \times CBDA)$  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.
- 11 The relative intensity of a characteristic ion in a sample analyte exceeded the acceptance with respect to the reference spectra, indicating interference.
- 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.
- 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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