



License #: 00000020LCVT89602592

Sample ID: 2310SMAZ0167.0465 Batch #: 10

Hemp THCa Flower

Batch #: 10

Strain: 10 Blue Gusher Parent Batch #: Sample Collected: Published: 10/30/2023 **Sample ID:** 2310SMAZ0167.0465

Amount Received: 5 g
Sample Type: Flower - Cured
Received: 10/18/2023

No Previev Available

COMPLIANCE FOR RETAIL

Regulated Analytes

Cannabinoid Profile (Q3)

Tested

Microbial Contaminants

Not Tested

Residual Solvents

Not Tested

Pesticides, Fungicides, and Growth Regulators

Pass

Mycotoxins

Not Tested

Heavy Metals

Pass

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

22.618%

18.045% Total THC

0.052%

Total CBD

ND

ND CBG

Total Cannabinoids (Q3)

Ahmed Munshi

Technical Laboratory Director

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Sample ID: 2310SMAZ0167.0465 Batch #: 10

Tested

Cannabinoid Profile

HPLC

Sample Prep

Batch Date: 10/18/2023 SOP: 418.AZ

Batch Number: 184

Sample Analysis

Date: 10/19/2023 SOP: 417.AZ - HPLC Sample Weight: 0.1079 g Volume: 40 mL

Analyte	LOD (mg/g)	LOQ (mg/g)	Dil.	Actual % (w/w)	mg/g	Qualifier
CBC	0.119	0.362	1	ND	ND	
CBD	0.119	0.362	1	ND	ND	
CBDA	0.119	0.362	1	0.059	0.593	
CBDV	0.119	0.362	1	ND	ND	
CBG	0.119	0.362	1	ND	ND	
CBGA	0.119	0.362	1	2.030	20.301	
CBN	0.119	0.362	1	ND	ND	
d8-THC	0.119	0.362	1	ND	ND	
d9-THC	0.119	0.362	1	0.241	2.412	
ТНСА	0.119	0.362	1	20.187	201.869	
THCV	0.119	0.362	1	ND	ND	

Cannabinoid Totals	Actual % (w/w)	mg/g	Qualifier		
Total THC	18.045	180.451			
Total CBD	0.052	0.520			
Total Cannabinoids	22.618	226.176	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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Sample ID: 2310SMAZ0167.0465 Batch #: 10

Heavy Metals

ICP-MS

Analyte

Pass

LOD (ppm)

Sample Prep

Batch Date: 10/19/2023 SOP: 428.AZ

Batch Number: 188

LOQ (ppm)

Sample Analysis

Date: 10/19/2023 SOP: 428.AZ - ICP-MS Sample Weight: 0.2230 g Volume: 6 mL

mit (ppm)	Results (ppm)	Qualifier

Arsenic	0.018	0.179	10	0.4	ND	
Cadmium	0.018	0.179	10	0.4	ND	
Lead	0.018	0.449	10	1	<loq< td=""><td></td></loq<>	
Mercury	0.018	0.090	10	0.2	ND	

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Pesticides, Fungicides, and Growth Regulators

LC-MS/MS Pass

Sample Prep

Batch Date: 10/18/2023 **SOP:** 432.AZ **Batch Number:** 178

Sample Analysis

Date: 10/19/2023 SOP: 424.AZ - LC-MS/MS Sample Weight: 0.5233 g Volume: 12.5 mL

Analyte	LOD / LOQ (ppm)	Dil.	Action Limit (ppm)	Results (ppm)	Qualifier	Analyte	LOD / LOQ (ppm)	Dil.	Action Limit (ppm)	Results (ppm)	Qualifier
Abamectin B1a	0.079 / 0.239	1	0.5	ND	I1	Hexythiazox	0.160 / 0.478	1	1	ND	M2
Acephate	0.064 / 0.191	1	0.4	ND		Imazalil	0.032 / 0.096	1	0.2	ND	
Acetamiprid	0.032 / 0.096	1	0.2	ND		Imidacloprid	0.064 / 0.191	1	0.4	ND	
Aldicarb	0.064 / 0.191	1	0.4	ND		Kresoxim-methyl	0.064 / 0.191	1	0.4	ND	
Azoxystrobin	0.032 / 0.096	1	0.2	ND		Malathion	0.032 / 0.096	1	0.2	ND	
Bifenazate	0.032 / 0.096	1	0.2	ND		Metalaxyl	0.032 / 0.096	1	0.2	ND	
Bifenthrin	0.032 / 0.096	1	0.2	ND	M2	Methiocarb	0.032 / 0.096	1	0.2	ND	M1
Boscalid	0.064 / 0.191	1	0.4	ND	M1	Methomyl	0.064 / 0.191	1	0.4	ND	
Carbaryl	0.032 / 0.096	1	0.2	ND		Myclobutanil	0.032 / 0.096	1	0.2	ND	M1
Carbofuran	0.032 / 0.096	1	0.2	ND		Naled	0.079 / 0.239	1	0.5	ND	
Chlorantraniliprole	0.032 / 0.096	1	0.2	ND	M1	Oxamyl	0.160 / 0.478	1	1	ND	
Chlorfenapyr	0.160 / 0.478	1	1	ND	I1, M1	Paclobutrazol	0.064 / 0.191	1	0.4	ND	M1
Chlorpyrifos	0.032 / 0.096	1	0.2	ND	M2	Permethrins	0.032 / 0.096	1	0.2	ND	I1, M2
Clofentezine	0.032 / 0.096	1	0.2	ND		Phosmet	0.032 / 0.096	1	0.2	ND	
Cyfluthrin	0.160 / 0.478	1	1	ND		Piperonyl Butoxide	0.318 / 0.955	1	2	ND	M2
Cypermethrin	0.160 / 0.478	1	1	ND	M2	Prallethrin	0.032 / 0.096	1	0.2	ND	
Daminozide	0.160 / 0.478	1	1	ND		Propiconazole	0.064 / 0.191	1	0.4	ND	
Diazinon	0.032 / 0.096	1	0.2	ND		Propoxur	0.032 / 0.096	1	0.2	ND	
Dichlorvos	0.016 / 0.048	1	0.1	ND		Pyrethrins	0.134 / 0.400	1	1	ND	I1, M1
Dimethoate	0.032 / 0.096	1	0.2	ND		Pyridaben	0.032 / 0.096	1	0.2	ND	M2
Ethoprophos	0.032 / 0.096	1	0.2	ND		Spinosad	0.032 / 0.096	1	0.2	ND	M2
Etofenprox	0.064 / 0.191	1	0.4	ND		Spiromesifen	0.032 / 0.096	1	0.2	ND	
Etoxazole	0.032 / 0.096	1	0.2	ND		Spirotetramat	0.032 / 0.096	1	0.2	ND	
Fenoxycarb	0.032 / 0.096	1	0.2	ND		Spiroxamine	0.064 / 0.191	1	0.4	ND	
Fenpyroximate	0.064 / 0.191	1	0.4	ND	M2	Tebuconazole	0.064 / 0.191	1	0.4	ND	
Fipronil	0.064 / 0.191	1	0.4	ND		Thiacloprid	0.032 / 0.096	1	0.2	ND	
Flonicamid	0.160 / 0.478	1	1	ND		Thiamethoxam	0.032 / 0.096	1	0.2	ND	
Fludioxonil	0.064 / 0.191	1	0.4	ND	M1	Trifloxystrobin	0.032 / 0.096	1	0.2	ND	M2

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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. 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, **B2** 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 L1 greater than the acceptance limits, but the sample's target analytes were not detected above the maximum allowable concentrations for the analytes in the The recovery from the matrix spike was high, but the recovery from the laboratory control sample was within acceptance criteria. M1 The recovery from the matrix spike was low, but the recovery from the laboratory control sample was within acceptance criteria. 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. 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. 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. 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. 02 The sample is heterogeneous, and sample homogeneity could not be readily achieved using routine laboratory practices. 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 Q3 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.

Notes:

V1

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maximum allowable for the analytes in the sample.

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The recovery from continuing calibration verification standards exceeded the acceptance limits, but the sample's target analytes were not detected above the