

TAPFIZZ.17.2024

Sample ID: 2404APO1724.8121
Strain: TAPFIZZ5:1V
Matrix: Ingestible
Type: Other
Source Batch #: HC-CPR1

Produced:
Collected: 04/23/2024 09:15 am
Received: 04/23/2024
Completed: 04/26/2024
Batch #: TAPFIZZ.17.2024
Harvest Date: 01/10/2023

Client
Michelle Mango
Lic. # 00000132ESFR75101840

Lot #:
Production Date: 04/22/2024
Production Method: Lecithin



Summary

Test	Date Tested	Result
Batch		
Cannabinoids	04/23/2024	Pass Complete
Microbials	04/26/2024	Pass

Cannabinoids by SOP-6

Complete

9.5711 mg/serving 95.7111 mg/container Total THC	<LOQ <LOQ Total CBD	9.9516 mg/serving 99.5157 mg/container Total Cannabinoids^(Q3)	NT Total Terpenes^(Q3)
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Analyte	LOD mg/container	LOQ mg/container	Result mg/container	Result mg/g	Q
THCa		1.8150	ND	ND	
Δ9-THC		1.8150	95.7111	3.1640	<div style="width: 100%; height: 10px; background-color: #0056b3;"></div>
Δ8-THC		1.8150	ND	ND	
THCV		1.8150	<LOQ	<LOQ	
CBDa		1.8150	ND	ND	
CBD		1.8150	<LOQ	<LOQ	
CBDVa		1.8150	ND	ND	
CBDV		1.8150	ND	ND	
CBN		1.8150	<LOQ	<LOQ	
CBGa		1.8150	ND	ND	
CBG		1.8150	3.8046	0.1258	<div style="width: 10%; height: 10px; background-color: #0056b3;"></div>
CBC		1.8150	<LOQ	<LOQ	
Total THC			95.7111	3.1640	
Total CBD			<LOQ	<LOQ	
Total			99.5157	3.2898	

Date Tested: 04/23/2024 07:00 am
10 servings per container.



Bryant Kearl
Lab Director
04/26/2024

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ARIZONA DEPARTMENT OF HEALTH SERVICES' WARNING:

Marijuana use can be addictive and can impair an individual's ability to drive a motor vehicle or operate heavy machinery. Marijuana smoke contains carcinogens and can lead to an increased risk for cancer, tachycardia, hypertension, heart attack, and lung infection. Marijuana use may affect the health of a pregnant woman and the unborn child. Using marijuana during pregnancy could cause birth defects or other health issues to your unborn child;

KEEP OUT OF REACH OF CHILDREN.

The product associated with the COA has been tested by Apollo Labs using validated state certified testing methodologies as required by Arizona state law. Values reported herein relate only to the specific sample of product submitted by Client for testing. Apollo Labs makes no claims as to the efficacy, safety or other risks associated with any detected or non-detected levels of any compounds reported herein. This Certificate shall not be reproduced except in full, without the written approval of Apollo Labs.

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Microbials

Pass

Analyte	Limit	Result	Status	Q
Salmonella SPP by QPCR: SOP-15	Detected/Not Detected in 1g	ND	Pass	
Aspergillus Flavus Aspergillus Fumigatus or Aspergillus Niger by QPCR: SOP-14	Detected/Not Detected in 1g	NR	NT	
Aspergillus Terreus by QPCR: SOP-14	Detected/Not Detected in 1g	NR	NT	

Analyte	LOQ	Limit	Result	Status	Q
E. Coli by traditional plating: SOP-13	CFU/g 10.0	CFU/g 10.0	CFU/g < 10 CFU/g	Pass	

Date Tested: 04/26/2024 12:00 am

Mycotoxins by SOP-22

Not Tested

Analyte	LOD	LOQ	Limit	Units	Status	Q
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Date Tested:

Heavy Metals by SOP-21

Not Tested

Analyte	LOD	LOQ	Limit	Units	Status	Q
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Date Tested:



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Qualifiers Definitions

Qualifier Notation	Qualifier Description
I1	The relative intensity of a characteristic ion in a sample analyte exceeded the acceptance criteria in subsection (L)(1) 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 in subsection (K)(2)(c), but the sample's target analytes were not detected above the maximum allowable concentrations in Table 3.1 for the analytes in the sample
M1	The recovery from the matrix spike in subsection (K)(4) was: a. High, but the recovery from the laboratory control sample in subsection (K)(2) was within acceptance criteria
M2	The recovery from the matrix spike in subsection (K)(4) was: b. Low, but the recovery from the laboratory control sample in subsection (K)(2) was within acceptance criteria
M3	The recovery from the matrix spike in subsection (K)(4) was: c. Unusable because the analyte concentration was disproportionate to the spike level, but the recovery from the laboratory control sample in subsection (K)(2) was within acceptance criteria
R1	The relative percent difference for the laboratory control sample and duplicate exceeded the limit in subsection (K)(3), but the recovery in subsection (K)(2) was within acceptance criteria
V1	The recovery from continuing calibration verification standards exceeded the acceptance limits in subsection (J)(1)(b), but the sample's target analytes were not detected above the maximum allowable concentrations in Table 3.1 for the analytes in the sample
Q2	The sample is heterogeneous, and sample homogeneity could not be readily achieved using routine laboratory practices - Used to denote that the sample as-received could not be fully pre-homogenized in packaging prior to microbiology analysis
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

Notes and Addenda:



Bryant Kearl
Lab Director
04/26/2024

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