

(Jeeter) JEETER CONCENTRATES APPLE FRITTER 1G CARTRIDGE

Sample ID: 2410EAZ0246.0973 Strain: APPLE FRITTER Matrix: Concentrates & Extracts Type: Vape Batch#: JJAZ-APPFRI-090824

Collected: 10/07/2024 Received: 10/07/2024 Completed: 10/09/2024 06:29 PM Sample Size: 17.5 g;

Harvest Date: 08/28/2024 Manufacture Date: 09/08/2024 External Lot ID#: Production Method: Butane

Client Jeeter

Lic. # 00000066DCBO00410690 2626 South Roosevelt Street, Tempe, AZ, 85282



Summarv

<u> </u>			
Test	Date Tested	Instr. Method	Result
Batch			Pass
Cannabinoids	10/08/2024	LC-UV VIS	Complete

Cannabinoids

Method: SOP AZ_M-CANNABINOIDS

86.153 %	0.153 0	%	91.465 %	
Total THC	Total CBD		Total Cannabinoids	
Analytes	LOQ	Result	Result	Q
	mg/g	%	mg/g	
THCA	0.769	ND	ND	Q3
Δ9 THC	0.769	86.153	861.53	Q3
Δ8 THC	0.769	ND	ND	Q3
THCVA	0.769	ND	ND	Q3
THCV	0.769	1.041	10.41	Q3
CBDA	0.769	ND	ND	Q3
CBD	0.769	0.153	1.53∎	Q3
CBN	0.769	0.273	2.73	Q3
CBGA	0.769	ND	ND	Q3
CBG	0.769	2.942	29.42	Q3
CBCA	0.769	ND	ND	Q3
CBC	0.769	0.903	9.03	Q3
Total THC	01100	86.153	861.53	Q3
Total CBD		0.153	1.53	Q3
Total Cannabinoids		91.465	914.65	Q3 Q3
Sum of Cannabinoids		91.465	914.65	Q3 Q3

Total THC = THCa * 0.877 + Δ9-THC; Total CBD = CBDa * 0.877 + CBD; Total Cannabinoids = (cannabinoid acid forms * 0.877) + cannabinoids; Sum of Cannabinoids = cannabinoid acid forms + cannabinoids; LOQ = Limit of Quantitation; NT = Not Tested; ND = Not Detected Moisture Method: SOP AZ_M-MOISTURE



Kevin Nolan Laboratory Technical Director | 10/09/2024





This product has been tested by Encore Labs Arizona using valid testing methodologies and a quality system as required by Arizona state law. Values reported relate only to the product tested. Encore 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 Encore Labs.

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Qualifier Legend

- The target analyte detected in the calibration blank required or the method blank is at or above the limit of quantitation, but the sample result for **B1** potency testing, is below the limit of quantitation.
- The target analyte detected in the calibration blank required or the method blank is at or above the limit of quantitation, but the sample result when **B2** testing for pesticides, fungicides, growth regulators, mycotoxins, heavy metals, or residual solvents, is below the maximum allowable concentration.
- **D1** The limit of quantitation and the sample results were adjusted to reflect sample dilution.
- The relative intensity of a characteristic ion in a sample analyte exceeded the acceptance with respect to the reference spectra, indicating 11 interference.
- When testing for pesticides, fungicides, herbicides, growth regulators, heavy metals, or residual solvents, the percent recovery of a laboratory control L1 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.
- The recovery from the matrix spike was high, but the recovery from the laboratory control sample was within acceptance criteria. M1
- M2 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 M3 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 M4 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 M5 sample.
- A description of the variance is described in the final report of testing according to R9-17- 404.06(B)(3)(d)(ii) N1
- Q1 Sample integrity was not maintained.
- Q2 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 Q3 requirements in R9-17-317.
- The relative percent difference for the laboratory control sample and duplicate exceeded the limit, but the recovery was within acceptance criteria. **R1**
- **R2** The relative percent difference for a sample and duplicate exceeded the limit.
- The recovery from initial or continuing calibration verification standards is greater than the acceptance limits, but the sample's target analytes were V1 not detected above the maximum allowable concentrations for the analytes in the sample.

Report Notes



Kevin Nolan Laboratory Technical Director | 10/09/2024





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