

Truffaloha - Live Rosin Oil

Sample ID: 2505EAZ0377.1722
Strain: Truffaloha
Matrix: Concentrates & Extracts
Type: Hash - Oil
Batch#: 5AC2Y

Collected: 05/20/2025 10:21 AM
Received: 05/20/2025
Completed: 05/22/2025
Sample Size: 63.70 g;

Harvest Date:
Manufacture Date:
External Lot ID#:
Production Method:

Client:
Canna Brands
Lic. # 0000156ESTDP70697204
6856 E Parkway Norte,
Mesa, AZ, 85212



Summary

Test	Date Tested	Instr. Method	Result
Batch			Pass
Cannabinoids	05/20/2025	LC-UV VIS	Complete
Terpenes	05/20/2025	GC-MS	Complete

Cannabinoids

Method: SOPAZ_M-CANNABINOIDS

77.281 %

Total THC

0.073 %

Total CBD

81.450 %

Total Cannabinoids ^{Q3}

Analytes	LOQ	Result	Result	Q
	mg/g	%	mg/g	
THCA	0.727	60.495	604.95	Q3
Δ9 THC	0.727	24.227	242.27	Q3
Δ8 THC	0.727	ND	ND	Q3
THCVA	0.727	0.378	3.78	Q3
THCV	0.727	0.161	1.61	Q3
CBDA	0.727	0.083	0.83	Q3
CBD	0.727	ND	ND	Q3
CBN	0.727	ND	ND	Q3
CBGA	0.727	1.915	19.15	Q3
CBG	0.727	0.592	5.92	Q3
CBCA	0.727	1.250	12.50	Q3
CBC	0.727	0.236	2.36	Q3
Total THC		77.281	772.81	Q3
Total CBD		0.073	0.73	Q3
Total Cannabinoids		81.450	814.50	Q3
Sum of Cannabinoids		89.337	893.37	Q3

Date Tested: 05/20/2025

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: SOPAZ_M-MOISTURE



Kevin Nolan
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Laboratory Technical Director | 05/22/2025

Firas Haddad
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Laboratory Manager | 05/22/2025



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Terpenes

Method: SOPAZ_M-TERPENES

Analytes	LOQ	Result	Result	Q
	mg/g	mg/g	%	
β-Caryophyllene	0.190	16.629	1.663	Q3
δ-Limonene	0.190	7.001	0.700	Q3
α-Humulene	0.190	6.624	0.662	Q3
β-Myrcene	0.190	6.371	0.637	Q3
Guaiol	0.951	4.916	0.492	Q3
α-Bisabolol	0.951	3.889	0.389	Q3
Linalool	0.190	2.139	0.214	Q3
β-Pinene	0.190	0.965	0.097	Q3
α-Pinene	0.190	0.908	0.091	Q3
trans-Nerolidol	0.228	0.776	0.078	Q3
Camphene	0.190	0.251	0.025	Q3
Terpinolene	0.190	<LOQ	<LOQ	Q3
δ-3-Carene	0.190	ND	ND	Q3
α-Terpinene	0.190	ND	ND	Q3
p-Cymene	0.190	ND	ND	Q3
Eucalyptol	0.190	ND	ND	Q3
cis-B-ocimene	0.190	ND	ND	Q3
trans-B-ocimene	0.190	ND	ND	Q3
γ-Terpinene	0.190	ND	ND	Q3
Isopulegol	0.951	ND	ND	Q3
Geraniol	0.951	ND	ND	Q3
cis-Nerolidol	0.380	ND	ND	Q3
Caryophyllene Oxide	0.951	ND	ND	Q3
Total		50.469	5.047	Q3

Date Tested: 05/20/2025

LOQ = Limit of Quantitation; NT = Not Tested; ND = Not Detected.

Primary Aromas

 Clove	 Citrusy	 Hops	 Musky	 Wood
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Qualifier Legend

- B1** *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 potency testing, is below the limit of quantitation.*
- B2** *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 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.*
- 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.*
- N1** *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 initial or continuing calibration verification standards 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.*

Report Notes




Kevin Nolan
Laboratory Technical Director | 05/22/2025


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