

Liv Life Infused Preroll Blueberry Muffin (LLIPBM9325)

Sample ID:
2509EAZ0641.3082

Strain: Blueberry Muffin
Matrix: Concentrates & Extracts

Type: Infused/Enhanced Preroll
Batch#: LLIPBM9325

Collected: 09/05/2025 10:30 AM

Received: 09/05/2025

Completed: 09/10/2025

Sample Size: 22.02 g;

Harvest Date: 12/30/2024

Manufacture Date: 09/03/2025

External Lot ID#: 42070925BLM ; 010125.ISO

Production Method: Butane

Client

Total Accountability Systems 1 Inc

Lic. # 000000009DCYP00763819, 00000110ESBL46708127

6287 E Copper Hill Dr,
Prescott Valley, AZ, 86314



Summary

Test	Date Tested	Instr. Method	Result
Batch			Pass
Cannabinoids	09/05/2025	LC-UV VIS	Complete
Microbial Impurities	09/08/2025	3M Plating & qPCR	Pass

Cannabinoids

Method: SOPAZ_M-CANNABINOIDS

45.028 %

Total THC

ND

Total CBD

45.634 %

Total Cannabinoids ^{Q3}

Analytes	LOQ	Result	Result	Q
	mg/g	%	mg/g	
THCA	0.755	50.585	505.85	
Δ9 THC	0.755	0.664	6.64	
Δ8 THC	0.755	ND	ND	
THCVA	0.755	0.146	1.46	
THCV	0.755	ND	ND	
CBDA	0.755	ND	ND	
CBD	0.755	ND	ND	
CBN	0.755	ND	ND	
CBGA	0.755	0.260	2.60	
CBG	0.755	ND	ND	
CBCA	0.755	0.286	2.86	
CBC	0.755	ND	ND	
Total THC		45.028	450.27	
Total CBD		ND	ND	
Total Cannabinoids		45.634	456.34	Q3
Sum of Cannabinoids		51.941	519.41	Q3

Date Tested: 09/05/2025

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 | 09/10/2025

Firas Haddad
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Laboratory Manager | 09/10/2025



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Microbial Impurities

Method: SOPAZ_M-ECOLI

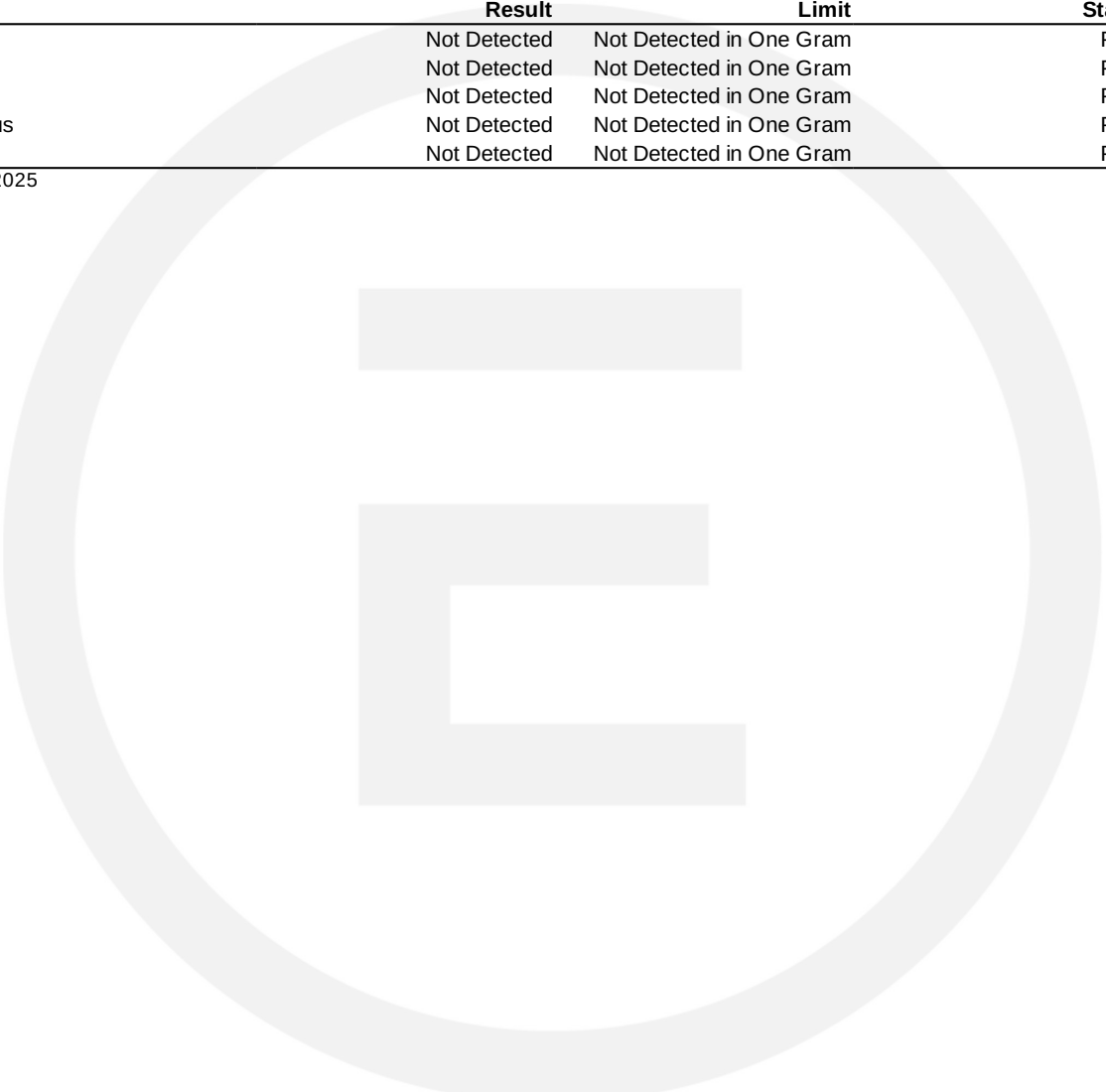
Analytes	Result	Limit	Status	Q
Escherichia coli	<10 CFU/g	100 CFU/g	Pass	

Date Tested: 09/08/2025

Method: SOPAZ_M-MICROBIALS

Analytes	Result	Limit	Status	Q
Salmonella spp	Not Detected	Not Detected in One Gram	Pass	
Aspergillus flavus	Not Detected	Not Detected in One Gram	Pass	
Aspergillus niger	Not Detected	Not Detected in One Gram	Pass	
Aspergillus fumigatus	Not Detected	Not Detected in One Gram	Pass	
Aspergillus terreus	Not Detected	Not Detected in One Gram	Pass	

Date Tested: 09/08/2025



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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



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