Encore Labs Arizona
16624 N 90th St, Suite 101
Scottsdale, AZ 85260
(626) 696-3086
https://encorelabs.com
Lic# 0000034LRCRF78097578

Certificate of Analysis

100MG Arnie

Sample ID: 2512EAZ0930.4436
Strain: Lime Glow
Matrix: Ingestible
Type: Beverage
Batch#: 095AR

Collected: 12/19/2025 12:50 PM
Received: 12/19/2025
Completed: 12/23/2025
Sample Size: 800.0 g;

Harvest Date:
Manufacture Date: 10/16/2025
External Lot ID#: LIOW.031424.FSE
Production Method:

Client
Canna Brands

Lic. # 00000133ESGJ79432018
1225 W Deer Valley
Phoenix, AZ

Summary

Test	Date Tested	Instr. Method	Result
Batch			Pass
Cannabinoids	12/19/2025	LC-UV VIS	Complete
Microbial Impurities	12/23/2025	3M Plating & qPCR	Pass

Cannabinoids

Method: SOP AZ_M-CANNABINOIDS

84.93 mg/unit	ND	84.93 mg/unit
Total THC	Total CBD	Total Cannabinoids Q3

Analytes	LOQ	Result	Result	Result	Q
	mg/g	%	mg/g	mg/unit	
THCA	0.020	ND	ND	ND	
Δ9 THC	0.020	0.012	0.12	84.93	
Δ8 THC	0.020	ND	ND	ND	
THCVA	0.020	ND	ND	ND	
THCV	0.020	ND	ND	ND	
CBDA	0.020	ND	ND	ND	
CBD	0.020	ND	ND	ND	
CBN	0.020	ND	ND	ND	
CBGA	0.020	ND	ND	ND	
CBG	0.020	ND	ND	ND	
CBCA	0.020	ND	ND	ND	
CBC	0.020	ND	ND	ND	
Total THC		**0.012**	**0.12**	**84.93**	
Total CBD		**ND**	**ND**	**ND**	
Total Cannabinoids		**0.012**	**0.12**	**84.93**	Q3
Sum of Cannabinoids		**0.012**	**0.12**	**84.93**	Q3

Date Tested: 12/19/2025
1 Unit = 712.75g;
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 | 12/23/2025

Firas Haddad
Laboratory Manager | 12/23/2025

ISO/IEC 17025:2017 Accredited
Accreditation # 122243

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Microbial Impurities

Method: SOP AZ_M-ECOLI

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

Date Tested: 12/23/2025

Method: SOP AZ_M-MICROBIALS

Analytes	Result	Limit	Status	Q
Salmonella spp	Not Detected	Not Detected in One Gram	Pass	

Date Tested: 12/23/2025

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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 between values obtained according to subsection N is more than 40%.*
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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