

Vape - Guava Lava 1000mg

Sample ID: 2508APO3622.18053

Strain: Guava Lava

Matrix: Concentrates & Extracts

Type: Vape

Source Batch #: The Desert Valley
Pharmacy Inc 00000060DCIS00424661;
Nature's Wonder Inc
00000035DCCB00049778

Collected: 08/18/2025 10:09 am

Received: 08/18/2025

Completed: 08/21/2025

Batch #: 08182550

Harvest Date: 05/29/2025

Client

Cure Injoy

Lic. # 00000060DCIS00424661

Lot #: 303NW0625

Production/Manufacture Date: 08/15/2025

Production/Manufacture Method: Alcohol



Summary

Test	Date Tested	Result
Batch		Pass
Cannabinoids	08/19/2025	Complete
Microbials	08/21/2025	Pass

Cannabinoids by SOP-6

Complete

84.4993 %	0.2005 %	88.7930 %	NT
Total THC	Total CBD	Total Cannabinoids ^(Q3)	Total Terpenes ^(Q3)

Analyte	LOD %	LOQ %	Result %	Result mg/g	Q
THCa		0.1000	<LOQ	<LOQ	
Δ9-THC		0.1000	84.4994	844.994	
Δ8-THC		0.1000	ND	ND	
THCV		0.1000	0.5315	5.315	
CBDa		0.1000	ND	ND	
CBD		0.1000	0.2006	2.006	
CBDVa		0.1000	ND	ND	
CBDV		0.1000	ND	ND	
CBN		0.1000	0.4035	4.035	
CBGa		0.1000	ND	ND	
CBG		0.1000	2.5013	25.013	
CBC		0.1000	0.6568	6.568	
Total THC			84.4993	844.9937	
Total CBD			0.2005	2.0057	
Total			88.7930	887.9304	

Date Tested: 08/19/2025 07:00 am



Anthony Settanni
Anthony Settanni
Lab Director
08/21/2025

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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	ND	Pass	
Aspergillus Terreus by QPCR: SOP-14	Detected/Not Detected in 1g	ND	Pass	

Analyte	LOQ	Limit	Result	Status	Q
E. Coli by traditional plating: SOP-13	CFU/g 10.0	CFU/g 100.0	CFU/g < 10 CFU/g	Pass	

Date Tested: 08/21/2025 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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Lab Director
08/21/2025

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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:



[Signature]

Bryant Kearl
Chief Scientific Officer
08/21/2025

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Production/Manufacture Method: Alcohol

Customer Supplied Information:

Source Batch: The Pharm / Sunday Goods Lic. # 00000099ESVM28064808



Bryant Kearl
Chief Scientific Officer
08/21/2025

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