Thanks For Choosing

License # 00000083CYO00463840

Product Information

PAGE 1: Product Type, Strain, Batch #, Weight, Potency, Harvest / Manufacture Date, Product Ingredients / Flower Nutrients, Production Chain, Extraction method (if applicable).

Page 2: Distribution Chain

PAGE 3: Product Warnings

PAGE 4 AND BEYOND: Product Testing

Product Type: Flower

Strain: AZ Super Sour OG

Batch #: 203961

THC : 23.03%
CBD: 0.00%

Total Cannabinoids: 26.00%

Harvest Date of Flower: 12/3/25

Production Chain:
Cultivated and Manufactured by Valley of the Sun Medical Dispensary License: 00000083DCYO00463840

Distribution Chain- Products Intended to be sold to the following Arizona Licensed Marijuana Dispensaries:

1. **Valley Of The Sun Medical Dispensary (License # 00000083CYO00463840)**

2. **All Greens Inc. (License # 00000011DCMZ00182361)**

PRODUCT WARNINGS

Valley of the Sun Medical Dispensary

Using marijuana during pregnancy could cause birth defects or other health issues to your unborn child

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. KEEP OUT OF REACH OF CHILDREN

Warning: There may be potential dangers to fetuses caused by smoking or ingesting marijuana while pregnant or to infants while breastfeeding

Warning: Use of marijuana during pregnancy may result in a risk being reported to the Department of Child Safety during pregnancy or at birth of the child by persons who are required to report

CERTIFICATE OF ANALYSIS

DATE ISSUED 01/21/2026 11:56 A.M. | CC ID: 2512C4L0027.2488

SAMPLE DETAILS

OVERALL BATCH RESULT: PASS

SAMPLE NAME: AZ Super Sour OG

Flower, Inhalable, Az Super Sour OG

CLIENT

Business Name: Valley Of the Sun Medical Dispensary INC.

License Number: 00000083dcyo00463840

Address: 16200 W Eddie Albert Way GOODYEAR AZ 85338

Scan QR code to verify authenticity of results.

SAMPLE DETAIL

Batch Number: 203961

Sample ID: 251217M042

Lot#:

Manufacture Date:

Harvest Date: 12/03/2025

Date Collected: 12/17/2025 11:55 a.m.

Date Received: 12/17/2025 2:16 p.m.

Batch Size:

Sample Size: 15.886 grams

Unit Mass:

Serving Size:

CANNABINOID ANALYSIS - SUMMARY

Sum of Cannabinoids: 26.00% (Q3)

Total Cannabinoids: 23.03% (Q3)

Total THC: 23.03%

Total CBD: ND

Sum of Cannabinoids = Δ^9-THC + THCa + CBD + CBDa + CBG + CBC + Δ^8-THC + CBN
Total Cannabinoids = (Δ^9-THC+0.877*THCa) + (CBD+0.877*CBDa) + CBG + CBC + Δ^8-THC + CBN
Total THC/CBD is calculated using the following formulas to take into account the loss of a carboxyl group during the decarboxylation step:
Total THC = Δ^9-THC + (THCa (0.877))
Total CBD = CBD + (CBDa (0.877))

*SAFETY ANALYSIS - SUMMARY

Pesticides: PASS **Heavy Metals:** PASS **Microbiology:** PASS **Microbiology (Plating):** PASS

***Amendment Note:** Positive detection of aspergillus contested at two separate laboratories, updated to non-detect.

These results relate only to the sample included on this report.
This report shall not be reproduced, except in full, without written approval of the laboratory.

Sample Certification: Testing results were obtained according to requirements in the quality assurance plan in R9-17-404.05, in the applicable standard operating procedure, and in R9-17-404.03 or R9-17-404.04. Results marked as 'Pass' or 'Fail' are done so in reference to R9-17: Arizona Administrative Code (A.A.C.) Title 9, Chapter 17.

Decision Rule: Statements of conformity (e.g. Pass/Fail) to specifications are made in this report without taking measurement uncertainty into account. Where statements of conformity are made in this report, the following decision rules are applied: PASS - Results within limits/specifications, FAIL - Results exceed limits/specifications.

References: limit of detection (LOD), limit of quantification (LOQ), not detected (ND), not tested (NT), µg/g = ppm, µg/kg = ppb, too numerous to count >250 cfu/plate (TNTC), colony-forming unit (cfu)

PJLA Testing
Accreditation #96186

Approved by: Mackenzie Whitman
Laboratory Director
Date: 01/21/2026

Amendment to Certificate of Analysis 251217M042-001

CANNABINOID TEST RESULTS - 12/18/2025

Tested by high-performance liquid chromatography with diode-array detection (HPLC-DAD). **Method:** (SOP-CHEM-003)

TOTAL CANNABINOIDS: 23.03% (Q3)
Total Cannabinoids (Total THC) + (Total CBD) + CBG + CBC + Δ^8-THC + CBN

TOTAL THC: 23.03%
Total THC (Δ^9-THC+0.877*THCa)

TOTAL CBD: ND
Total CBD (CBD+0.877*CBDa)

COMPOUND	LOD/LOQ (mg/g)	QUALIFIERS	RESULT (mg/g)	RESULT (%)
THCa	*0.5 / 2.6*		241.2	24.12
Δ^9-THC	*0.5 / 2.6*		18.8	1.88
CBDa	*0.4 / 2.6*		<LOQ	<LOQ
CBG	*0.3 / 2.6*		<LOQ	<LOQ
CBC	*0.5 / 2.6*		<LOQ	<LOQ
Δ^8-THC	*0.6 / 2.6*		ND	ND
CBD	*0.6 / 2.6*		ND	ND
CBN	*0.4 / 2.6*		ND	ND
SUM OF CANNABINOIDS (Q3)			260.0 mg/g	26.00%

PESTICIDE TEST RESULTS - 12/22/2025 PASS

Pesticide and plant growth regulator analysis utilizing high-performance liquid chromatography-mass spectrometry (HPLC-MS/MS). **Method:** (SOP-CHEM-006)

COMPOUND	LOD/LOQ (µg/g)	ACTION LIMIT (µg/g)	QUALIFIERS	RESULT (µg/g)	RESULT
Abamectin	*0.085 / 0.111*	*0.5*	*V1*	ND	PASS
Acephate	*0.022 / 0.092*	*0.4*	*L1,V1*	ND	PASS
Acetamiprid	*0.017 / 0.046*	*0.2*		ND	PASS
Aldicarb	*0.045 / 0.092*	*0.4*	*L1*	ND	PASS
Azoxystrobin	*0.012 / 0.046*	*0.2*	*L1,V1*	ND	PASS
Bifenazate	*0.023 / 0.046*	*0.2*	*L1,V1*	ND	PASS
Bifenthrin	*0.017 / 0.046*	*0.2*		ND	PASS
Boscalid	*0.068 / 0.092*	*0.4*	*V1*	ND	PASS
Carbaryl	*0.022 / 0.046*	*0.2*	*L1*	ND	PASS
Carbofuran	*0.012 / 0.046*	*0.2*	*L1*	ND	PASS
Chlorantraniliprole	*0.028 / 0.046*	*0.2*	*L1,V1*	ND	PASS
Chlorfenapyr	*0.334 / 0.461*	*1*	*R1,I1*	ND	PASS
Chlorpyrifos	*0.026 / 0.046*	*0.2*		ND	PASS
Clofentezine	*0.012 / 0.046*	*0.2*		ND	PASS
Cyfluthrin	*0.235 / 0.461*	*1*	*I1*	ND	PASS
Cypermethrin	*0.095 / 0.231*	*1*	*L1,V1*	ND	PASS
Daminozide	*0.063 / 0.461*	*1*	*L1,V1*	ND	PASS

PESTICIDE TEST RESULTS - 12/22/2025 *continued*

COMPOUND	LOD/LOQ (µg/g)	ACTION LIMIT (µg/g)	QUALIFIERS	RESULT (µg/g)	RESULT
Diazinon	*0.013 / 0.046*	*0.2*	*L1*	ND	PASS
Dichlorvos (DDVP)	*0.012 / 0.046*	*0.1*	*L1,I1,V1*	ND	PASS
Dimethoate	*0.014 / 0.046*	*0.2*	*L1,V1*	ND	PASS
Ethoprophos	*0.015 / 0.046*	*0.2*	*L1,V1*	ND	PASS
Etofenprox	*0.028 / 0.092*	*0.4*		ND	PASS
Etoxazole	*0.015 / 0.046*	*0.2*	*L1*	ND	PASS
Fenoxycarb	*0.015 / 0.046*	*0.2*		ND	PASS
Fenpyroximate	*0.036 / 0.092*	*0.4*	*L1,V1*	ND	PASS
Fipronil	*0.062 / 0.092*	*0.4*		ND	PASS
Flonicamid	*0.065 / 0.231*	*1*	*L1,V1*	ND	PASS
Fludioxonil	*0.046 / 0.092*	*0.4*		ND	PASS
Hexythiazox	*0.075 / 0.231*	*1*		ND	PASS
Imazalil	*0.019 / 0.046*	*0.2*		ND	PASS
Imidacloprid	*0.039 / 0.092*	*0.4*	*V1*	ND	PASS
Kresoxim-methyl	*0.040 / 0.092*	*0.4*		ND	PASS
Malathion	*0.049 / 0.046*	*0.2*	*L1,V1*	ND	PASS
Metalaxyl	*0.015 / 0.046*	*0.2*	*L1*	ND	PASS
Methiocarb	*0.037 / 0.046*	*0.2*	*L1,V1*	ND	PASS
Methomyl	*0.024 / 0.092*	*0.4*		ND	PASS
Myclobutanil	*0.026 / 0.046*	*0.2*	*I1*	ND	PASS
Naled	*0.026 / 0.115*	*0.5*	*L1,V1*	ND	PASS
Oxamyl	*0.057 / 0.231*	*1*		ND	PASS
Paclobutrazol	*0.033 / 0.092*	*0.4*	*L1,V1*	ND	PASS
Permethrins	*0.024 / 0.046*	*0.2*	*V1*	ND	PASS
Phosmet	*0.015 / 0.046*	*0.2*	*L1,I1,V1*	ND	PASS
Piperonyl Butoxide	*0.142 / 0.461*	*2*	*L1,V1*	ND	PASS
Prallethrin	*0.012 / 0.046*	*0.2*	*L1,V1*	ND	PASS
Propiconazole	*0.067 / 0.092*	*0.4*	*L1,V1*	ND	PASS
Propoxur	*0.019 / 0.046*	*0.2*		ND	PASS
Pyrethrins	*0.049 / 0.129*	*1*		ND	PASS
Pyridaben	*0.011 / 0.046*	*0.2*	*L1,V1*	ND	PASS
Spinosad	*0.017 / 0.036*	*0.2*		ND	PASS
Spiromesifen	*0.017 / 0.046*	*0.2*		ND	PASS
Spirotetramat	*0.033 / 0.046*	*0.2*	*L1,V1*	ND	PASS
Spiroxamine	*0.022 / 0.092*	*0.4*		ND	PASS
Tebuconazole	*0.045 / 0.092*	*0.4*	*L1,V1*	ND	PASS
Thiacloprid	*0.017 / 0.046*	*0.2*		ND	PASS
Thiamethoxam	*0.015 / 0.046*	*0.2*	*L1,V1*	ND	PASS
Trifloxystrobin	*0.017 / 0.046*	*0.2*	*V1*	ND	PASS

HEAVY METALS TEST RESULTS - 12/22/2025 PASS

Heavy metal analysis utilizing inductively coupled plasma-mass spectrometry (ICP-MS). **Method:** (SOP-CHEM-008)

COMPOUND	LOD/LOQ (µg/g)	ACTION LIMIT (µg/g)	QUALIFIERS	RESULT (µg/g)	RESULT
Arsenic	*0.01 / 0.10*	*0.4*		<LOQ	PASS
Cadmium	*0.01 / 0.10*	*0.4*		ND	PASS
Lead	*0.02 / 0.40*	*1*		<LOQ	PASS
Mercury	*0.01 / 0.04*	*0.2*		<LOQ	PASS

*MICROBIOLOGY TEST RESULTS - 01/21/2026 PASS

Analysis conducted by polymerase chain reaction (PCR) and fluorescence detection of microbiological contaminants. **Method:** (SOP-MICRO-017)

COMPOUND	QUALIFIERS	RESULT	RESULT
Aspergillus flavus		Not Detected in 1 gram	PASS
Aspergillus fumigatus		Not Detected in 1 gram	PASS
Aspergillus niger		Not Detected in 1 gram	PASS
Aspergillus terreus		Not Detected in 1 gram	PASS
Salmonella spp.		Not Detected in 1 gram	PASS

***Amendment Note:** Positive detection of aspergillus contested at two separate laboratories, updated to non-detect.

*MICROBIOLOGY TEST RESULTS - 01/21/2026 PASS

Analysis conducted by 3M™ Petrifilm™. **Method:** (SOP-MICRO-010)

COMPOUND	LOQ (cfu/g)	ACTION LIMIT (cfu/g)	QUALIFIERS	RESULT (cfu/g)	RESULT
Escherichia coli	*10*	*100*		<10	PASS

***Amendment Note:** Positive detection of aspergillus contested at two separate laboratories, updated to non-detect.

Notes and Defnitions

Item	Definition
L1	When testing for pesticides, fungicides, growth regulators, mycotoxins, heavy metals, or residual solvents, the percent recovery of a laboratory controlsample 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.
R1	The relative percent difference for the laboratory control sample and duplicate exceeded the limit, but the recovery was within acceptance criteria.
I1	The relative intensity of a characteristic ion in a sample analyte exceeded the acceptance criteria with respect to the reference spectra, indicating interference.
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.
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. Testing result is not accredited under ISO 17025.
Notes	

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. KEEP OUT OF REACH OF CHILDREN. Using Marijuana during pregnancy could cause birth defects or other health issues to your unborn child.

Valley Of the Sun Medical Dispensary INC.
16200 W Eddie Albert Way
GOODYEAR
AZ
85338
License #: 00000083dcyo00463840
Sample ID: 2601SMAZ0026.0084
Batch #: 203961

Certificate: 19962

SMITHERS

CERTIFICATE OF ANALYSIS

License #: 00000020LCVT89602592

AZ Super Sour OG

Batch #: 203961
Strain: AZ Super Sour OG
Parent Batch #:
Production Method: Indoor
Harvest Date: 12/03/2025
Received: 01/07/2026

Sample ID: 2601SMAZ0026.0084
Amount Received: 6.4 g
Sample Type: Flower - Cured
Sample Collected: 01/07/2026 16:21:00
Manufacture Date:
Published: 01/10/2026

COMPLIANCE FOR RETAIL

Regulated Analytes

Cannabinoid Profile (Q3) Not Tested	Microbial Contaminants **Pass**	Residual Solvents Not Tested
Pesticides, Fungicides, and Growth Regulators Not Tested	Mycotoxins Not Tested	Heavy Metals Not Tested

Additional Analytes (Not Regulated)

Terpenes Total (Q3) Not Tested	Moisture Analysis (Q3) Not Tested	Water Activity (Q3) Not Tested
Filth & Foreign (Q3) Not Tested	Homogeneity (Q3) Not Tested	Additional Microbial Contaminants (Q3) Not Tested

NT Total THC
NT Total CBD
NT CBN
NT CBG
NT Total Cannabinoids (Q3)

Ahmed Munshi

Technical Laboratory Director

Smithers CTS Arizona LLC
734 W Highland Avenue, 2nd Floor
Phoenix, AZ 85013
(602) 806-6930

PJLA Testing
Accreditation #: 103104

The product associated with this COA has been tested by Smithers CTS Arizona LLC, using validated state certified testing methodologies as required by Arizona state law.
Testing results were obtained according to Smithers' quality assurance plan and requirements found in R9-17-404.03 and R9-17-404.04.
This COA is governed by the terms and conditions listed on: https://www.smithers.com/arizona-terms-conditions

Valley Of the Sun Medical Dispensary INC.
16200 W Eddie Albert Way
GOODYEAR
AZ
85338
License #: 00000083dcyo00463840
Sample ID: 2601SMAZ0026.0084
Batch #: 203961

Certificate: 19962

SMITHERS

CERTIFICATE OF ANALYSIS

License #: 00000020LCVT89602592

Microbial Analysis

Pass

Sample Prep

Batch Date: 01/08/2026
SOP: 406.AZ
Batch Number: 4916
Test ID: 109661

Sample Analysis

Date: 01/09/2026
SOP: 406.AZ - qPCR (MG)
Sample Weight: 1.020 g

Analyte	Allowable Criteria	Actual Result	Pass/Fail	Qualifier
Aspergillus flavus	Not Detected in One Gram	Not Detected in One Gram	Pass	
Aspergillus fumigatus	Not Detected in One Gram	Not Detected in One Gram	Pass	
Aspergillus niger	Not Detected in One Gram	Not Detected in One Gram	Pass	
Aspergillus terreus	Not Detected in One Gram	Not Detected in One Gram	Pass	

Ahmed Munshi

Technical Laboratory Director

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Phoenix, AZ 85013
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Valley Of the Sun Medical Dispensary INC.
16200 W Eddie Albert Way
GOODYEAR
AZ
85338
License #: 00000083dcyo00463840
Sample ID: 2601SMAZ0026.0084
Batch #: 203961

Certificate: 19962

SMITHERS

CERTIFICATE OF ANALYSIS

License #: 00000020LCVT89602592

Qualifier Legend

B1 *The target analyte detected in the calibration 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, or the method blank is at or above the limit of quantitation, but the sample result when testing for pesticides, fungicides, herbicides, growth regulators, heavy metals, or residual solvents, is below the maximum allowable concentration for the analyte.*

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.*

M6 *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 continuing calibration verification standards exceeded the acceptance limits, but the sample's target analytes were not detected above the maximum allowable for the analytes in the sample.*

Cultivated By:
Manufactured By:
Disclaimer: Using marijuana during pregnancy could cause birth defects or other health issues to your unborn child.

Ahmed Munshi

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Valley Of the Sun Medical Dispensary INC.
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SMITHERS

CERTIFICATE OF ANALYSIS

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Notes:

Ahmed Munshi

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PJLA Testing
Accreditation #: 103104

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

AZ Super Sour OG

Sample ID: 2601EAZ0015.0042
Strain: AZ Super Sour OG
Matrix: Plant
Type: Flower - Cured
Batch#: 203961

Collected: 01/12/2026 02:09 PM
Received: 01/12/2026
Completed: 01/14/2026
Sample Size: 7.57 g;

Harvest Date: 12/03/2025
Manufacture Date:
External Lot ID#:
Production Method: Indoor

Client
Valley Of the Sun Medical Dispensary INC.

Lic. # 00000083dcyo00463840
16200 W Eddie Albert Way,
GOODYEAR, AZ, 85338

Summary

Test	Date Tested	Instr. Method	Result
Batch			Pass
Microbial Impurities	01/14/2026	3M Plating & qPCR	Pass

Microbial Impurities

Method: SOP AZ_M-MICROBIALS

Analytes	Result	Limit	Status	Q
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: 01/14/2026

Firas Haddad
Laboratory Manager | 01/14/2026

ISO/IEC 17025:2017 Accredited
Accreditation # 122243

AZ Super Sour OG

Sample ID: 2601EAZ0015.0042
Strain: AZ Super Sour OG
Matrix: Plant
Type: Flower - Cured
Batch#: 203961

Collected: 01/12/2026 02:09 PM
Received: 01/12/2026
Completed: 01/14/2026
Sample Size: 7.57 g;

Harvest Date: 12/03/2025
Manufacture Date:
External Lot ID#:
Production Method: Indoor

Client
Valley Of the Sun Medical Dispensary INC.

Lic. # 00000083dcyo00463840
16200 W Eddie Albert Way,
GOODYEAR, AZ, 85338

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

Firas Haddad
Laboratory Manager | 01/14/2026

ISO/IEC 17025:2017 Accredited
Accreditation # 122243