TRANSPARENT LABS

RSP Productions: Asylum

2985 W. Whitton Ave.
Phoenix, AZ 85017
lodellrichardson@gmail.com
(702) 338-7088
Lic. #00000070DCBD00783295
Harvest Dates: 04/30/2024

Isis x ONYCD Flower

Plant, Flower - Cured
Dispensary License #: ; Manufacturing License #: ; Cultivation License #:

Sample: 2405TLL0157.0774

Strain: Isis x ONYCD
Parent Batch #: ; Batch#: 20240430R9ISO; Batch Size: 15 g
Sample Received: 05/06/2024; Report Created: 05/10/2024; Expires: 05/10/2025
Manufacturing Date:
Sampling: ; Environment:

Safety

Pass	Pass	Pass
Pesticides	Microbials	Metals

Cannabinoids
TPL_Potency_01

24.19%	ND	28.42%
Total THC	Total CBD	Total Cannabinoids Q3

Analyte	LOQ	Mass	Mass	Qualifier
	%	%	mg/g	
THCa	0.10	25.87	258.7	
Δ9-THC	0.10	1.51	15.1	
Δ8-THC	0.10	ND	ND	
THCV	0.10	ND	ND	
CBDa	0.10	ND	ND	
CBD	0.10	ND	ND	
CBDV	0.10	ND	ND	
CBN	0.10	ND	ND	
CBGa	0.10	0.91	9.1	
CBG	0.10	0.14	1.4	
CBC	0.10	ND	ND	
Total		**28.42**	**284.2**	

Total THC = THCa * 0.877 + Δ9-THC
Total CBD = CBDa * 0.877 + CBD
Instrument: HPLC-DAD: ; Method: TPL_Potency_01

Terpenes
TPL_Terpenes_01

Cinnamon	Hops	Earthy

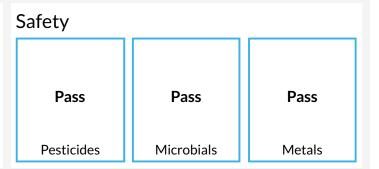| Analyte	LOQ	Mass	Mass	Qualifier
	%	%	mg/g	
β-Caryophyllene	0.1500	1.500	Q3
β-Myrcene	0.1000	1.000	Q3
Ocimene	0.1000	1.000	Q3
β-Pinene	0.0900	0.900	Q3
δ-Limonene	0.0900	0.900	Q3
α-Humulene	0.0700	0.700	Q3
γ-Terpinene	0.0700	0.700	Q3
Terpinolene	0.0700	0.700	Q3
trans-Nerolidol	0.0600	0.600	Q3
α-Pinene	0.0300	0.300	Q3
α-Bisabolol	0.0200	0.200	Q3
Camphene	0.0100	0.100	Q3
Linalool	0.0100	0.100	Q3
3-Carene	<	<	Q3
α-Terpinene	<	<	Q3
Caryophyllene Oxide	<	<	Q3
cis-Nerolidol	<	<	Q3
Eucalyptol	<	<	Q3
Geraniol	<	<	Q3
Guaiol	<	<	Q3
Isopulegol	<	<	Q3
p-Cymene	<	<	Q3
Total	**0.8700**	**8.700**	

Instrument: GCMS; Method: TPL_Terp_01
Notes:

1721 E McDowell Road
Phoenix, AZ
(602) 368-4233
transparentlabsaz.com
Lic# 0000029LRCXG19240160

Brian DiMarco
Laboratory Director

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coa.support@confidentlims.com
(866) 506-5866
www.confidentlims.com

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Certificate of Analysis

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Pesticides TPL_Pesticides_01   Pass

Analyte	LOQ PPM	Limit PPM	Mass PPM	Status	Qualifier	Analyte	LOQ PPM	Limit PPM	Mass PPM	Status	Qualifier
Abamectin	0.24	0.50	ND	Pass	V1	Hexythiazox	0.48	1.00	ND	Pass	L1, V1
Acephate	0.19	0.40	ND	Pass		Imazalil	0.10	0.20	ND	Pass	M2
Acetamiprid	0.10	0.20	ND	Pass	V1	Imidacloprid	0.19	0.40	ND	Pass	
Aldicarb	0.19	0.40	ND	Pass		Kresoxim Methyl	0.19	0.40	ND	Pass	
Azoxystrobin	0.10	0.20	ND	Pass		Malathion	0.10	0.20	ND	Pass	
Bifenazate	0.10	0.20	ND	Pass	M1	Metalaxyl	0.10	0.20	ND	Pass	
Bifenthrin	0.10	0.20	ND	Pass	L1, V1	Methiocarb	0.10	0.20	ND	Pass	M2
Boscalid	0.19	0.40	ND	Pass		Methomyl	0.19	0.40	ND	Pass	
Carbaryl	0.10	0.20	ND	Pass		Myclobutanil	0.10	0.20	ND	Pass	L1, V1
Carbofuran	0.10	0.20	ND	Pass		Naled	0.24	0.50	ND	Pass	
Chlorantraniliprole	0.10	0.20	ND	Pass		Oxamyl	0.48	1.00	ND	Pass	
Chlorfenapyr	0.48	1.00	ND	Pass	M2, V1	Paclobutrazol	0.19	0.40	ND	Pass	
Chlorpyrifos	0.10	0.20	ND	Pass	M1	Permethrin	0.10	0.20	ND	Pass	M2, V1
Clofentezine	0.10	0.20	ND	Pass	V1	Phosmet	0.10	0.20	ND	Pass	
Cyfluthrin	0.48	1.00	ND	Pass	V1	Piperonyl Butoxide	0.96	2.00	<LOQ	Pass	
Cypermethrin	0.48	1.00	ND	Pass	M1, V1	Prallethrin	0.10	0.20	ND	Pass	M1, L1
Daminozide	0.48	1.00	ND	Pass	M1	Propiconazole	0.19	0.40	ND	Pass	M2
Diazinon	0.10	0.20	ND	Pass		Propoxur	0.10	0.20	ND	Pass	
Dichlorvos	0.05	0.10	ND	Pass		Pyrethrins	0.48	1.00	ND	Pass	L1
Dimethoate	0.10	0.20	ND	Pass		Pyridaben	0.10	0.20	ND	Pass	V1
Ethoprophos	0.10	0.20	ND	Pass		Spinosad	0.10	0.20	ND	Pass	
Etofenprox	0.19	0.40	ND	Pass		Spiromesifen	0.10	0.20	ND	Pass	
Etoxazole	0.10	0.20	ND	Pass		Spirotetramat	0.10	0.20	ND	Pass	
Fenoxycarb	0.10	0.20	ND	Pass		Spiroxamine	0.19	0.40	ND	Pass	
Fenpyroximate	0.19	0.40	ND	Pass		Tebuconazole	0.19	0.40	ND	Pass	
Fipronil	0.19	0.40	ND	Pass		Thiacloprid	0.10	0.20	ND	Pass	
Flonicamid	0.48	1.00	ND	Pass		Thiamethoxam	0.10	0.20	ND	Pass	
Fludioxonil	0.19	0.40	ND	Pass	M2	Trifloxystrobin	0.10	0.20	ND	Pass	

Instrument: LC-QQQ ; Method: TPL_Pesticides_01

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Heavy Metals — Pass

Analyte	LOQ	Limit	Mass	Status	Qualifier
	PPB	PPB	PPB		
Arsenic	200.0	400.0	<LOQ	Pass	V1
Cadmium	200.0	400.0	<LOQ	Pass	
Lead	500.0	1000.0	<LOQ	Pass	
Mercury	100.0	200.0	<LOQ	Pass	L1 R1

Microbials — Pass

Analyte	LOQ	Limit	Result	Status	Qualifier
	CFU/g	CFU/g	CFU/g		
E. Coli	10	100	<10	Pass	

Analyte	Limit	Result	Status	Qualifier
Salmonella	Detectable in 1g	Not Detected	Pass	
Aspergillus	Detectable in 1g	Not Detected	Pass	
Aspergillus fumigatus	Detectable in 1g	Not Detected	Pass	
Aspergillus niger	Detectable in 1g	Not Detected	Pass	
Aspergillus flavus	Detectable in 1g	Not Detected	Pass	
Aspergillus terreus	Detectable in 1g	Not Detected	Pass	

LOQ=Limit of Quantitation. The reported result is based on a simple weight with the applicable moisture content for that sample. Unless otherwise stated, all quality control samples performed within specifications established by the Laboratory. Instrument: ICPMS; Method: AOAC 2021.03

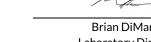Instrument: qPCR/Plating; AOAC Methods 082102, 022202 and 2018.13

B1 = Target analyte detected in calibration blank was above LOQ but the concentration of cannabinoid was blow LOQ,

B2 = Target analyte detected in calibration blank was above LOQ but was 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 criteria with respect to the reference spectra, indicating interference,

L1 = The percent recovery of a laboratory control sample is greater than the acceptance limits in A.A.C 17 R9-17-404.03(K)(2)(C), but the sample's target analytes were not detected above the maximum allowed concentration,

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,

R1 = The relative percent difference for the laboratory control sample and duplicate exceeded the limit in A.A.C 17 R9-17-404.03(K)(3), but the recover in subsection A.A.C 17 R9-17-404.03 (K)(2) was within accepted criteria,

R2 = The relative percent difference for a sample and duplicated exceeded the limit in subsection A.A.C 17 R9-17-404.03 (O)

Q1 = Sample integrity was not maintained,

Q2 = The sample is heterogenous 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

V1 = The recovery from continuing calibration verification standards exceeded the acceptance limits denoted in A.C.C 17 R9-17-403.03(J)(1)(b), but the sample's target analytes were not detected above the maximum allowable concentrations for the analytes in the sample.

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