

## Total Health & Wellness dba True Harvest

Sample: 2405TLL0165.0807

Phoenix, AZ 85043  
jpastor@trueharvestco.com

Strain: Cadillac Rainbows x All Dressed  
Parent Batch #: ; Batch#: CRAD240509; Batch Size: 17 g  
Sample Received: 05/10/2024; Report Created: 05/17/2024; Expires: 05/17/2025  
Manufacturing Date:  
Sampling: ; Environment:

Lic. #00000100DCWU00857159  
Harvest Dates:

## Cadillac Rainbows x All Dressed

Concentrates & Extracts, Infused/Enhanced Preroll, Extraction Method: Ice/Water  
Dispensary License #: ; Manufacturing License #: ; Cultivation License #:



## Safety

Pass Pesticides	Pass Microbials	Pass Mycotoxins
Pass Solvents	Pass Metals	Not Tested Foreign Matter

## Cannabinoids

TPL\_Potency\_01

35.97% Total THC	<LOQ Total CBD	41.98% Total Cannabinoids Q3
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Analyte	LOQ	Mass	Mass	Qualifier
	%	%	mg/g	
THCa	0.10	37.47	374.7	
Δ9-THC	0.10	3.11	31.1	
Δ8-THC	0.10	ND	ND	
THCV	0.10	ND	ND	
CBDa	0.10	<LOQ	<LOQ	
CBD	0.10	ND	ND	
CBDV	0.10	ND	ND	
CBN	0.10	ND	ND	
CBGa	0.10	1.19	11.9	
CBG	0.10	0.21	2.1	
CBC	0.10	ND	ND	
<b>Total</b>		<b>41.98</b>	<b>419.8</b>	

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	 Earthy	 Lemon
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Analyte	LOQ	Mass	Mass	Qualifier
	%	%	mg/g	
β-Caryophyllene		0.2830	2.830	Q3
Ocimene		0.2220	2.220	Q3
δ-Limonene		0.2040	2.040	Q3
Guaiol		0.2000	2.000	Q3
β-Myrcene		0.1740	1.740	Q3
β-Pinene		0.1520	1.520	Q3
Terpinolene		0.1490	1.490	Q3
α-Humulene		0.1280	1.280	Q3
Eucalyptol		0.0880	0.880	Q3
trans-Nerolidol		0.0610	0.610	Q3
Linalool		0.0570	0.570	Q3
α-Pinene		0.0150	0.150	Q3
α-Bisabolol		0.0130	0.130	Q3
3-Carene		<	<	Q3
α-Terpinene		<	<	Q3
Camphene		<	<	Q3
Caryophyllene Oxide		<	<	Q3
cis-Nerolidol		<	<	Q3
γ-Terpinene		<	<	Q3
Geraniol		<	<	Q3
Isopulegol		<	<	Q3
p-Cymene		<	<	Q3
<b>Total</b>		<b>1.7460</b>	<b>17.460</b>	

Instrument: GCMS; Method: TPL\_Terp\_01  
Notes:

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## Pesticides TPL\_Pesticides\_01

Pass

Analyte	LOQ	Limit	Mass	Status	Qualifier	Analyte	LOQ	Limit	Mass	Status	Qualifier
	PPM	PPM	PPM				PPM	PPM	PPM		
Abamectin	0.24	0.50	ND	Pass	V1 L1	Hexythiazox	0.48	1.00	ND	Pass	
Acephate	0.19	0.40	ND	Pass		Imazalil	0.10	0.20	ND	Pass	
Acetamiprid	0.10	0.20	ND	Pass		Imidacloprid	0.19	0.40	ND	Pass	M1
Aldicarb	0.19	0.40	ND	Pass		Kresoxim	0.19	0.40	ND	Pass	
Azoxystrobin	0.10	0.20	ND	Pass		Methyl					
Bifenazate	0.10	0.20	ND	Pass	V1 M1	Malathion	0.10	0.20	ND	Pass	L1
Bifenthrin	0.10	0.20	ND	Pass	L1 M2	Metalaxyl	0.10	0.20	ND	Pass	
Boscalid	0.19	0.40	ND	Pass		Methiocarb	0.10	0.20	ND	Pass	
Carbaryl	0.10	0.20	ND	Pass		Methomyl	0.19	0.40	ND	Pass	
Carbofuran	0.10	0.20	ND	Pass		Myclobutanil	0.10	0.20	ND	Pass	
Chlorantraniliprole	0.10	0.20	ND	Pass		Naled	0.24	0.50	ND	Pass	
Chlorfenapyr	0.48	1.00	ND	Pass	L1 V1	Oxamyl	0.48	1.00	ND	Pass	
Chlorpyrifos	0.10	0.20	ND	Pass	M1	Paclobotrazol	0.19	0.40	ND	Pass	L1
Clofentezine	0.10	0.20	ND	Pass	M2	Permethrin	0.10	0.20	ND	Pass	L1 M2
Cyfluthrin	0.48	1.00	ND	Pass	M1 L1	Phosmet	0.10	0.20	ND	Pass	
Cypermethrin	0.48	1.00	ND	Pass	M1 L1	Piperonyl	0.96	2.00	ND	Pass	
Daminozide	0.48	1.00	ND	Pass	M1 L1	Butoxide					
Diazinon	0.10	0.20	ND	Pass	V1	Prallethrin	0.10	0.20	ND	Pass	L1 M1
Dichlorvos	0.05	0.10	ND	Pass		Propiconazole	0.19	0.40	ND	Pass	
Dimethoate	0.10	0.20	ND	Pass		Propoxur	0.10	0.20	ND	Pass	
Ethoprophos	0.10	0.20	ND	Pass		Pyrethrins	0.48	1.00	ND	Pass	
Etofenprox	0.19	0.40	ND	Pass	L1 M2	Pyridaben	0.10	0.20	ND	Pass	
Etoazole	0.10	0.20	ND	Pass	L1	Spinosad	0.10	0.20	ND	Pass	I1
Fenoxycarb	0.10	0.20	ND	Pass		Spiromesifen	0.10	0.20	ND	Pass	
Fenproximate	0.19	0.40	ND	Pass		Spirotetramat	0.10	0.20	ND	Pass	M1
Fipronil	0.19	0.40	ND	Pass	M1	Spiroxamine	0.19	0.40	ND	Pass	
Flonicamid	0.48	1.00	ND	Pass		Tebuconazole	0.19	0.40	ND	Pass	
Fludioxonil	0.19	0.40	ND	Pass		Thiacloprid	0.10	0.20	ND	Pass	
						Thiamethoxam	0.10	0.20	ND	Pass	
						Trifloxystrobin	0.10	0.20	ND	Pass	

Instrument: LC-QQ ; 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	ND	Pass	V1
Cadmium	200.0	400.0	<LOQ	Pass	V1
Lead	500.0	1000.0	<LOQ	Pass	V1
Mercury	100.0	200.0	<LOQ	Pass	V1

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

### Residual Solvents Pass

Analyte	LOQ	Limit	Mass	Status	Qualifier
	PPM	PPM	PPM		
Acetone	490.7	1000.0	ND	Pass	
Acetonitrile	201.2	410.0	ND	Pass	
Benzene	1.0	2.0	ND	Pass	
Butanes	613.4	5000.0	ND	Pass	
Chloroform	29.4	60.0	ND	Pass	
Dichloromethane	294.4	600.0	ND	Pass	
Ethanol	2453.4	5000.0	ND	Pass	
Ethyl-Acetate	2453.4	5000.0	ND	Pass	
Ethyl-Ether	2453.4	5000.0	ND	Pass	
Heptane	2453.4	5000.0	ND	Pass	
Hexanes	142.3	290.0	ND	Pass	
Isopropyl-Acetate	2453.4	5000.0	ND	Pass	
Methanol	1472.0	3000.0	ND	Pass	
Pentanes	142.3	5000.0	ND	Pass	
2-Propanol	2453.4	5000.0	ND	Pass	
Toluene	436.7	890.0	ND	Pass	
Xylenes	98.1	2170.0	ND	Pass	

Performed by GCMS-HS SOP-004. Methods used per AZDHS R9-17-404.03 and the solvent limits set by AZDHS R9-17 Table 3.1. AZDHS approved method for residual solvents by GCMS-HS for all listed analytes. Subcontracted through DVT Registration Certificate Identification Number : 0000031LRCHX78341676

### 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	

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

### Mycotoxins Pass

Analyte	LOQ	Limit	Mass	Status	Qualifier
	PPB	PPB	PPB		
B1	8.1	20.0	ND	Pass	
B2	8.1	20.0	ND	Pass	
G1	8.1	20.0	ND	Pass	
G2	8.1	20.0	ND	Pass	
Ochratoxin A	8.1	20.0	ND	Pass	I1
Total Aflatoxins	8.1	20.0	ND	Pass	

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

V1 = The recovery from continuing calibration verification standards exceeded the acceptance limits denoted in A.A.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.