

Certificate of Analysis

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Metals

Total Health & Wellness dba True Harvest

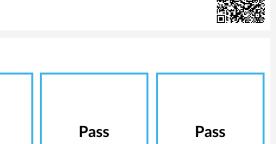
Sample: 2401TLL0023.0141

Phoenix, AZ 85043 jpastor@trueharvestco.com

Strain: Sour Dubb Parent Batch #:; Batch#: R5SD1228; Batch Size: 20 g Sample Received: 01/23/2024; Report Created: 01/26/2024; Expires: 01/26/2025 Manufacturing Date: Sampling: ; Environment:

Lic. #00000100DCWU00857159 Harvest Dates: Sour Dubb

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





Cannabinoids

TPL_Potency_01		_		
28.90%	<loq< th=""><th></th><th>33.56%</th><th>6</th></loq<>		33.56%	6
Total THC	Total CBD		Total Cannabinoids Q3	
Analyte	LOQ	Mass	Mass	Qualifier
	%	%	mg/g	
THCa	0.20	32.45	324.5	
∆9-THC	0.20	0.45	4.5	
∆8-THC	0.20	ND	ND	
THCV	0.20	ND	ND	
CBDa	0.20	<loq< td=""><td><loq< td=""><td></td></loq<></td></loq<>	<loq< td=""><td></td></loq<>	
CBD	0.20	ND	ND	
CBDV	0.20	ND	ND	
CBN	0.20	ND	ND	
CBGa	0.20	0.66	6.6	
CBG	0.20	<loq< th=""><th><loq< th=""><th></th></loq<></th></loq<>	<loq< th=""><th></th></loq<>	
CBC	0.20	ND	ND	
Total		33.56	335.6	

Total THC = THCa * 0.877 + Δ 9-THC Total CBD = CBDa * 0.877 + CBD Instrument: HPLC-DAD: ; Method: TPL_Potency_01



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α-Humulene 0.83 β-Caryophyllene 0.73 β-Myrcene 0.69 Linalool 0.47 trans-Nerolidol 0.41 β-Pinene 0.28 α-Bisabolol 0.19 Guaiol 0.16 Terpinolene 0.13 0.11 α-Pinene 0.11 y-Terpinene 0.07 0.7 Eucalyptol 0.04 Camphene 0.4 3-Carene α-Terpinene Caryophyllene Oxide cis-Nerolidol Geraniol Isopulegol Ocimene p-Cymene Total 5.08 50.8

Instrument: GCMS; Method: TPL_Terp_01

Notes:

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Brian DiMarco Laboratory Director

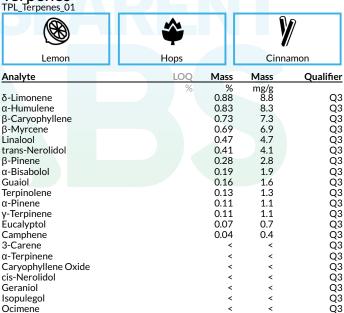
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Pass

Pesticides

Safety



Microbials



Harvest Dates:

Certificate of Analysis

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Sample: 2401TLL0023.0141

Total Health & Wellness dba True Harvest

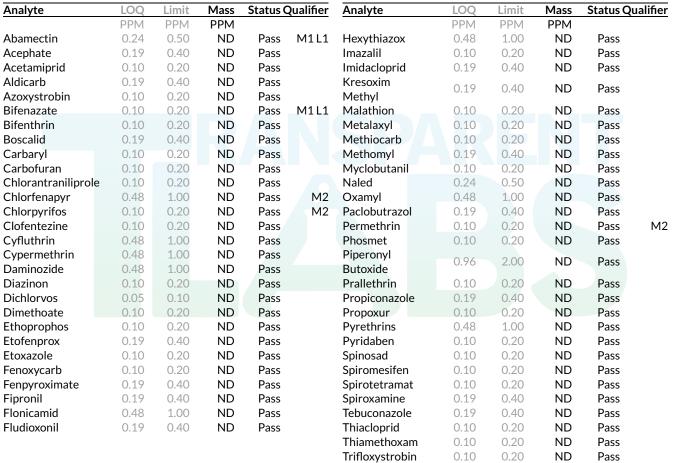
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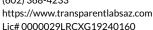
Pesticides TPL Pesticides 01



Instrument: LC-QQQ; Method: TPL_Pesticides_01



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Laboratory Director

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Pass



Certificate of Analysis

Pass

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Strain: Sour Dubb

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Sample: 2401TLL0023.0141

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Heavy Metals

Analyte	LOQ	Limit	Mass	Status	Qualifier
	PPB	PPB	PPB		
Arsenic	200.0	400.0	ND	Pass	
Cadmium	200.0	400.0	<loq< th=""><th>Pass</th><th></th></loq<>	Pass	
Lead	500.0	1000.0	<loq< th=""><th>Pass</th><th></th></loq<>	Pass	
Mercury	100.0	200.0	<loq< th=""><th>Pass</th><th></th></loq<>	Pass	

Microbials				Pass
Analyte	LOQ	Limit	Result	StatusQualifier
	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: ICPMS; Method: AOAC 2021.03

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



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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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