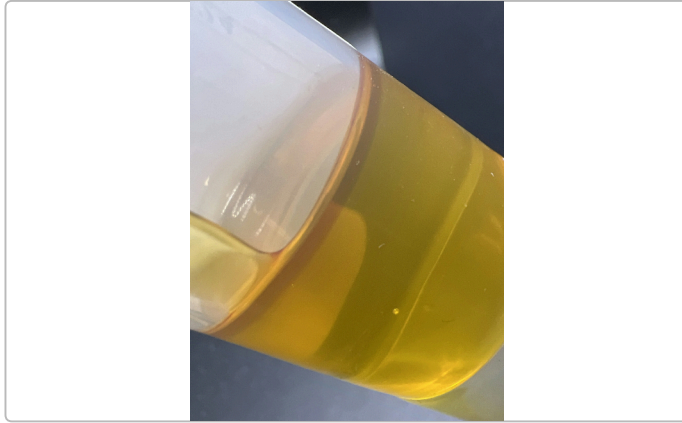


## DIME Berry White

Sample ID: 2403APO1036.4935  
Strain: Berry White  
Matrix: Concentrates & Extracts  
Type: Distillate  
Source Batch #: W C3L.FLV022524

Produced:  
Collected: 03/11/2024 12:04 pm  
Received: 03/11/2024  
Completed: 03/14/2024  
Batch #: BW0308  
Harvest Date: 01/10/2024

Client  
**Dime Industries**  
Lic. # 00000075ESJK64208740  
  
Lot #: W C3L.FLV022524  
Production Date:  
Production Method: Alcohol



### Summary

Test	Date Tested	Result
Batch		
Cannabinoids	03/12/2024	Complete
Microbials	03/14/2024	Pass

### Cannabinoids by SOP-6

Complete

<b>94.2634%</b>	<b>0.5316%</b>	<b>98.3627%</b>	<b>NT</b>
Total THC	Total CBD	Total Cannabinoids <sup>(Q3)</sup>	Total Terpenes <sup>(Q3)</sup>

Analyte	LOD	LOQ	Result	Result	Q
	%	%	%	mg/g	
THCa		0.1000	ND	ND	
Δ9-THC		0.1000	94.2634	942.634	<div style="width: 94.2634%;"></div>
Δ8-THC		0.1000	ND	ND	
THCV		0.1000	0.4589	4.589	<div style="width: 0.4589%;"></div>
CBDa		0.1000	ND	ND	
CBD		0.1000	0.5316	5.316	<div style="width: 0.5316%;"></div>
CBDVa		0.1000	ND	ND	
CBDV		0.1000	ND	ND	
CBN		0.1000	0.5361	5.361	<div style="width: 0.5361%;"></div>
CBGa		0.1000	ND	ND	
CBG		0.1000	1.5754	15.754	<div style="width: 1.5754%;"></div>
CBC		0.1000	0.9972	9.972	<div style="width: 0.9972%;"></div>
<b>Total THC</b>			<b>94.2634</b>	<b>942.6340</b>	
<b>Total CBD</b>			<b>0.5316</b>	<b>5.3160</b>	
<b>Total</b>			<b>98.3627</b>	<b>983.627</b>	

Date Tested: 03/12/2024 07:00 am



Bryant Kearl  
Lab Director  
03/14/2024

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## DIME Berry White

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Strain: Berry White  
Matrix: Concentrates & Extracts  
Type: Distillate  
Source Batch #: WC3L.FLV022524

Produced:  
Collected: 03/11/2024 12:04 pm  
Received: 03/11/2024  
Completed: 03/14/2024  
Batch #: BW0308  
Harvest Date: 01/10/2024

Client  
**Dime Industries**  
Lic. # 00000075ESJK64208740  
  
Lot #: WC3L.FLV022524  
Production Date:  
Production Method: Alcohol

### 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: 03/14/2024 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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Client  
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Lot #: WC3L.FLV022524  
Production Date:  
Production Method: Alcohol

**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:**



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Lab Director  
03/14/2024

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## WC3L Distillate

Sample ID: 2402APO0842.3914  
Strain: Distillate  
Matrix: Concentrates & Extracts  
Type: Distillate  
Source Batch #:

Produced:  
Collected: 02/26/2024 02:28 pm  
Received: 02/26/2024  
Completed: 02/29/2024  
Batch #: WC3L.FLV022524  
Harvest Date: 01/10/2024

Client  
**White Mountain Health Center - FLAV**  
Lic. # 00000083ESGB09219996

Lot #:  
Production Date: 02/13/2024  
Production Method: Butane



### Summary

Test	Date Tested	Result
Batch		Pass
Cannabinoids	02/27/2024	Complete
Residual Solvents	02/28/2024	Pass
Microbials	02/29/2024	Pass
Mycotoxins	02/27/2024	Pass
Pesticides	02/27/2024	Pass
Heavy Metals	02/27/2024	Pass

### Cannabinoids

Complete

<b>94.2539%</b>	<b>0.1847%</b>	<b>99.6227%</b>	<b>NT</b>
Total THC	Total CBD	Total Cannabinoids <sup>(Q3)</sup>	Total Terpenes <sup>(Q3)</sup>

Analyte	LOD	LOQ	Result	Result	Q
	%	%	%	mg/g	
THCa		0.1000	ND	ND	
Δ9-THC		0.1000	94.2539	942.539	
Δ8-THC		0.1000	ND	ND	
THCV		0.1000	0.3956	3.956	
CBDa		0.1000	ND	ND	
CBD		0.1000	0.1847	1.847	
CBDVa		0.1000	ND	ND	
CBDV		0.1000	ND	ND	
CBN		0.1000	0.5835	5.835	
CBGa		0.1000	ND	ND	
CBG		0.1000	2.3718	23.718	
CBC		0.1000	1.8331	18.331	
<b>Total THC</b>			<b>94.2539</b>	<b>942.5390</b>	
<b>Total CBD</b>			<b>0.1847</b>	<b>1.8470</b>	
<b>Total</b>			<b>99.6227</b>	<b>996.227</b>	

Date Tested: 02/27/2024 07:00 am



Bryant Kearl  
Lab Director  
02/29/2024

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Strain: Distillate  
Matrix: Concentrates & Extracts  
Type: Distillate  
Source Batch #:

Produced:  
Collected: 02/26/2024 02:28 pm  
Received: 02/26/2024  
Completed: 02/29/2024  
Batch #: WC3L.FLV022524  
Harvest Date: 01/10/2024

Client  
**White Mountain Health Center - FLAV**  
Lic. # 00000083ESGB09219996  
  
Lot #:  
Production Date: 02/13/2024  
Production Method: Butane

## Pesticides

Pass

Analyte	LOQ	Limit	Mass	Q	Status	Analyte	LOQ	Limit	Mass	Q	Status
	PPM	PPM	PPM				PPM	PPM	PPM		
Abamectin	0.2500	0.5000	ND		Pass	Hexythiazox	0.5000	1.0000	ND		Pass
Acephate	0.2000	0.4000	ND		Pass	Imazalil	0.1000	0.2000	ND		Pass
Acetamiprid	0.1000	0.2000	ND		Pass	Imidacloprid	0.2000	0.4000	ND		Pass
Aldicarb	0.2000	0.4000	ND		Pass	Kresoxim Methyl	0.2000	0.4000	ND		Pass
Azoxystrobin	0.1000	0.2000	ND		Pass	Malathion	0.1000	0.2000	ND		Pass
Bifenazate	0.1000	0.2000	0.1336	M1	Pass	Metalaxyl	0.1000	0.2000	ND		Pass
Bifenthrin	0.1000	0.2000	ND		Pass	Methiocarb	0.1000	0.2000	ND		Pass
Boscalid	0.2000	0.4000	ND		Pass	Methomyl	0.2000	0.4000	ND		Pass
Carbaryl	0.1000	0.2000	ND		Pass	Myclobutanil	0.1000	0.2000	ND		Pass
Carbofuran	0.1000	0.2000	ND		Pass	Naled	0.2500	0.5000	ND		Pass
Chlorantraniliprole	0.1000	0.2000	ND		Pass	Oxamyl	0.5000	1.0000	ND		Pass
Chlorfenapyr	0.5000	1.0000	ND		Pass	Pacllobutrazol	0.2000	0.4000	ND		Pass
Chlorpyrifos	0.1000	0.2000	ND		Pass	Permethrins	0.1000	0.2000	ND	M2	Pass
Clofentezine	0.1000	0.2000	ND		Pass	Phosmet	0.1000	0.2000	ND		Pass
Cyfluthrin	0.5000	1.0000	ND		Pass	Piperonyl	1.0000	2.0000	ND		Pass
Cypermethrin	0.5000	1.0000	ND		Pass	Butoxide	0.1000	0.2000	ND	M2	Pass
Daminozide	0.5000	1.0000	ND		Pass	Prallethrin	0.2000	0.4000	ND		Pass
Diazinon	0.1000	0.2000	ND		Pass	Propiconazole	0.2000	0.4000	ND		Pass
Dichlorvos	0.0500	0.1000	ND		Pass	Propoxur	0.1000	0.2000	ND		Pass
Dimethoate	0.1000	0.2000	ND		Pass	Pyrethrins	0.5000	1.0000	ND		Pass
Ethoprophos	0.1000	0.2000	ND		Pass	Pyridaben	0.1000	0.2000	ND		Pass
Etofenprox	0.2000	0.4000	ND	M2	Pass	Spinosad	0.1000	0.2000	ND	M2	Pass
Etoxazole	0.1000	0.2000	ND		Pass	Spiromesifen	0.1000	0.2000	ND		Pass
Fenoxycarb	0.1000	0.2000	ND		Pass	Spirotetramat	0.1000	0.2000	ND		Pass
Fenpyroximate	0.2000	0.4000	ND		Pass	Spiroxamine	0.2000	0.4000	ND		Pass
Fipronil	0.2000	0.4000	ND		Pass	Tebuconazole	0.2000	0.4000	ND	M2	Pass
Flonicamid	0.5000	1.0000	ND		Pass	Thiacloprid	0.1000	0.2000	ND		Pass
Fludioxonil	0.2000	0.4000	ND		Pass	Thiamethoxam	0.1000	0.2000	ND		Pass
						Trifloxystrobin	0.1000	0.2000	ND		Pass

Date Tested: 02/27/2024 07:00 am



Bryant Kearl  
Lab Director  
02/29/2024

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## WC3L Distillate

Sample ID: 2402APO0842.3914  
Strain: Distillate  
Matrix: Concentrates & Extracts  
Type: Distillate  
Source Batch #:

Produced:  
Collected: 02/26/2024 02:28 pm  
Received: 02/26/2024  
Completed: 02/29/2024  
Batch #: WC3L.FLV022524  
Harvest Date: 01/10/2024

Client  
**White Mountain Health Center - FLAV**  
Lic. # 00000083ESGB09219996

Lot #:  
Production Date: 02/13/2024  
Production Method: Butane

## Microbials

Pass

Analyte	Limit	Result	Status	Q
Salmonella SPP	Detected/Not Detected in 1g	ND	Pass	
Aspergillus Flavus Aspergillus Fumigatus or Aspergillus Niger	Detected/Not Detected in 1g	ND	Pass	
Aspergillus terreus	Detected/Not Detected in 1g	ND	Pass	

Analyte	LOQ	Limit	Result	Status	Q
E. Coli	CFU/g 10.0	CFU/g 100.0	CFU/g < 10 CFU/g	Pass	

Date Tested: 02/29/2024 12:00 am

## Mycotoxins

Pass

Analyte	LOD	LOQ	Limit	Units	Status	Q
	µg/kg	µg/kg	µg/kg	µg/kg		
B1	5	10	20	ND	Pass	
B2	5	10	20	ND	Pass	
G1	5	10	20	ND	Pass	
G2	5	10	20	ND	Pass	R1
Total Aflatoxins	5	10	20	ND	Pass	R1
Ochratoxin A	5	10	20	ND	Pass	

Date Tested: 02/27/2024 07:00 am

## Heavy Metals

Pass

Analyte	LOD	LOQ	Limit	Units	Status	Q
	PPM	PPM	PPM	PPM		
Arsenic	0.0660	0.1330	0.4000	ND	Pass	V1
Cadmium	0.0660	0.1330	0.4000	ND	Pass	V1
Lead	0.1660	0.3330	1.0000	ND	Pass	V1
Mercury	0.0330	0.0660	0.2000	ND	Pass	

Date Tested: 02/27/2024 07:00 am



Bryant Kearl  
Lab Director  
02/29/2024

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Harvest Date: 01/10/2024

Client  
**White Mountain Health Center - FLAV**  
Lic. # 00000083ESGB09219996

Lot #:  
Production Date: 02/13/2024  
Production Method: Butane

## Residual Solvents

Analyte	LOQ	Limit	Mass	Status	Q
	PPM	PPM	PPM		Pass
Acetone	381.0000	1000.0000	ND	Pass	
Acetonitrile	154.0000	410.0000	ND	Pass	
Benzene	1.0000	2.0000	ND	Pass	
Butanes	1914.0000	5000.0000	ND	Pass	
Chloroform	24.0000	60.0000	ND	Pass	
Dichloromethane	231.0000	600.0000	ND	Pass	
Ethanol	1910.0000	5000.0000	ND	Pass	
Ethyl-Acetate	1907.0000	5000.0000	ND	Pass	
Ethyl-Ether	1901.0000	5000.0000	ND	Pass	
n-Heptane	1892.0000	5000.0000	ND	Pass	
Hexanes	115.0000	290.0000	ND	Pass	
Isopropanol	1915.0000	5000.0000	ND	Pass	
Isopropyl-Acetate	1908.0000	5000.0000	ND	Pass	
Methanol	1141.0000	3000.0000	ND	Pass	
Pentane	1923.0000	5000.0000	ND	Pass	
Toluene	343.0000	890.0000	ND	Pass	
Xylenes + Ethyl Benzene	841.0000	2170.0000	ND	Pass	

Date Tested: 02/28/2024 07:00 am



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Lab Director  
02/29/2024

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Client  
**White Mountain Health Center - FLAV**  
Lic. # 00000083ESGB09219996

Lot #:  
Production Date: 02/13/2024  
Production Method: Butane

**Terpenes**

Analyte	LOQ	Mass	Mass	Q	Analyte	LOQ	Mass	Mass	Q
---------	-----	------	------	---	---------	-----	------	------	---



**Primary Aromas**

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Date Tested:



Bryant Kearl  
Lab Director  
02/29/2024

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Matrix: Concentrates & Extracts  
Type: Distillate  
Source Batch #:

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Client  
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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:**



Bryant Kearl  
Lab Director  
02/29/2024

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



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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.