

Genesis Biocenticals, LLC

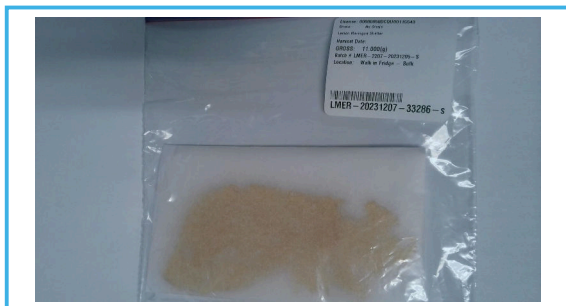
1120 W Watkins St
Phoenix, AZ 85007
shonae.j@genbioaz.com
(847) 682-4899
Lic. #00000058DCQU00115543
Harvest Dates:

Sample: 2312TLL0063.0171

Strain: Lemon Meringue
Parent Batch #: ; Batch#: LMER-2207-20231205-S; Batch Size: 11 g
Sample Received: 12/11/2023; Report Created: 12/21/2023; Expires: 12/21/2024
Manufacturing Date:
Sampling: ; Environment:

Lemon Meringue Shatter

Concentrates & Extracts, Shatter, Extraction Method: Butane
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

80.72% Total THC	0.18% Total CBD	NT Moisture
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Analyte	LOQ	Mass	Mass	Qualifier
	%	%	mg/g	
THCa	0.10	90.08	900.8	
Δ9-THC	0.10	1.72	17.2	
Δ8-THC	0.10	NR	NR	
THCV	0.10	NR	NR	
CBDa	0.10	0.20	2.0	
CBD	0.10	NR	NR	
CBDV	0.10	NR	NR	
CBN	0.10	NR	NR	
CBGa	0.10	7.17	71.7	
CBG	0.10	0.37	3.7	
CBC	0.10	NR	NR	
Total		99.55	995.5	

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

Terpenes

TPL_Terpenes_01

 Hops	 Cinnamon	 Lavender
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Analyte	LOQ	Mass	Mass	Qualifier
	%	%	mg/g	
α-Humulene		1.9280	19.280	Q3
β-Caryophyllene		1.6490	16.490	Q3
Linalool		1.1710	11.710	Q3
trans-Nerolidol		0.9110	9.110	Q3
Guaiol		0.5820	5.820	Q3
β-Myrcene		0.3780	3.780	Q3
β-Pinene		0.3750	3.750	Q3
Terpinolene		0.1840	1.840	Q3
α-Pinene		0.1630	1.630	Q3
3-Carene		0.1620	1.620	Q3
Ocimene		0.1610	1.610	Q3
γ-Terpinene		0.1100	1.100	Q3
Camphene		0.0540	0.540	Q3
α-Bisabolol		0.0460	0.460	Q3
Isopulegol		0.0230	0.230	Q3
Caryophyllene Oxide		NR	NR	
δ-Limonene		NR	NR	
Eucalyptol		NR	NR	
Nerolidol		NR	NR	
p-Cymene		NR	NR	
α-Terpinene		<	<	Q3
cis-Nerolidol		<	<	Q3
Geraniol		<	<	Q3
Total		7.8970	78.970	

Instrument: GCMS; Method: TPL_Terp_01
Notes:

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Residual Solvents

Pass

Analyte	LOQ	Limit	Mass	Status	Qualifier
	PPM	PPM	PPM		
Acetone	196.0	1000.0	<LOQ	Pass	
Acetonitrile	80.0	410.0	ND	Pass	
Benzene	0.4	2.0	ND	Pass	
n-Butane			ND	Tested	
Isobutane			ND	Tested	
Butanes	490.0	5000.0	<LOQ	Pass	
Chloroform	12.0	60.0	ND	Pass	
Dichloromethane	118.0	600.0	ND	Pass	
Ethanol	980.0	5000.0	ND	Pass	
Ethyl-Acetate	980.0	5000.0	ND	Pass	
Ethyl-Ether	980.0	5000.0	ND	Pass	
Heptane	980.0	5000.0	ND	Pass	
n-Hexane			ND	Tested	
2-Methyl-Pentane			ND	Tested	
3-Methyl-Pentane			ND	Tested	
2,2-Dimethyl-Butane			ND	Tested	
2,3-Dimethyl-Butane			ND	Tested	
Hexanes	142.0	290.0	ND	Pass	
Isopropyl-Acetate	980.0	5000.0	ND	Pass	
Methanol	588.0	3000.0	ND	Pass	
n-Pentane			ND	Tested	
Isopentane			ND	Tested	
Neopentane			ND	Tested	
Pentanes	980.0	5000.0	ND	Pass	
2-Propanol	980.0	5000.0	<LOQ	Pass	
Propane			ND	Tested	
Toluene	175.0	890.0	ND	Pass	
1,2-Dimethyl-Benzene			ND	Tested	
1,3-Dimethyl-Benzene			ND	Tested	
1,4-Dimethyl-Benzene			ND	Tested	
Ethyl-Benzene			ND	Tested	
Xylenes	851.0	2170.0	ND	Pass	

Heavy Metals

Pass

Analyte	LOQ	Limit	Mass	Status	Qualifier
	PPB	PPB	PPB		
Arsenic	200.0	400.0	ND	Pass	
Cadmium	200.0	400.0	ND	Pass	
Lead	500.0	1000.0	ND	Pass	
Mercury	100.0	200.0	ND	Pass	M2

Mycotoxins

Pass

Analyte	LOQ	Limit	Mass	Status	Qualifier
	PPB	PPB	PPB		
B1	10	0	ND	Pass	
B2	10	0	ND	Pass	
G1	10	0	ND	Pass	
G2	5	0	ND	Pass	I1,V1
Ochratoxin A	10	20	ND	Pass	I1,M1
Total Aflatoxins	10	20	ND	Pass	V1

Instrument: HS-GCMS

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Laboratory Director

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The product associated with this COA has been tested by Transparent Labs using state validated testing methods, as required by The State of Arizona. Measurement uncertainty and decision rule information is available upon request. The test results on this COA are only valid for the sample submitted by the client and are not valid for samples or batches not mentioned on this Certificate of Analysis. Transparent 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 COA shall not be reproduced except in full, except without the written approval of Transparent Labs. The required tests and associated limit values are referenced from The required tests and testing limits used within this COA conform to those specified in A.R.S Title 36, Chapter 28.2 and A.A.C Title 9 Chapter 17 Supp. 22-3. Using Marijuana during pregnancy could cause birth defects or other health issues to your unborn child.



Certificate of Analysis

Powered by Confident Cannabis
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Microbials

Pass

Analyte	LOQ	Limit	Result	Status	Qualifier
E. Coli	CFU/g 10	CFU/g 100	CFU/g <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: Plating/qPRC ; Method: AOAC 82102 and 022202



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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		Hexythiazox	0.48	1.00	ND	Pass	M2
Acephate	0.19	0.40	ND	Pass		Imazalil	0.10	0.20	ND	Pass	I1
Acetamiprid	0.10	0.20	ND	Pass		Imidacloprid	0.19	0.40	ND	Pass	
Aldicarb	0.19	0.40	ND	Pass		Kresoxim	0.19	0.40	ND	Pass	M2
Azoxystrobin	0.10	0.20	ND	Pass		Methyl					
Bifenazate	0.10	0.20	ND	Pass	L1	Malathion	0.10	0.20	ND	Pass	
Bifenthrin	0.10	0.20	ND	Pass	L1,V1	Metalaxyl	0.10	0.20	ND	Pass	
Boscalid	0.19	0.40	ND	Pass	M2	Methiocarb	0.10	0.20	ND	Pass	L1
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	
Chlorantranilprole	0.10	0.20	ND	Pass	L1	Naled	0.24	0.50	ND	Pass	
Chlorfenapyr	0.48	1.00	ND	Pass	I1	Oxamyl	0.48	1.00	ND	Pass	
Chlorpyrifos	0.10	0.20	ND	Pass		Paclobutrazol	0.19	0.40	ND	Pass	L1,V1
Clofentezine	0.10	0.20	ND	Pass	M2	Permethrin	0.10	0.20	ND	Pass	
Cyfluthrin	0.48	1.00	ND	Pass	M2	Phosmet	0.10	0.20	ND	Pass	
Cypermethrin	0.48	1.00	ND	Pass	M2, V1	Piperonyl	0.95	2.00	ND	Pass	
Daminozide	0.48	1.00	ND	Pass		Butoxide					
Diazinon	0.10	0.20	ND	Pass		Prallethrin	0.10	0.20	ND	Pass	
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	L1	Pyrethrins	0.48	1.00	ND	Pass	
Etofenprox	0.19	0.40	ND	Pass	L1,V1	Pyridaben	0.10	0.20	ND	Pass	M2
Etoxazole	0.10	0.20	ND	Pass		Spinosad	0.10	0.20	ND	Pass	M2
Fenoxycarb	0.10	0.20	ND	Pass		Spiromesifen	0.10	0.20	ND	Pass	
Fenpyroximate	0.19	0.40	ND	Pass	M2	Spirotetramat	0.10	0.20	ND	Pass	
Fipronil	0.19	0.40	ND	Pass	I1,R1	Spiroxamine	0.19	0.40	ND	Pass	
Flonicamid	0.48	1.00	ND	Pass		Tebuconazole	0.19	0.40	ND	Pass	L1,V1
Fludioxonil	0.19	0.40	ND	Pass	M2	Thiacloprid	0.10	0.20	ND	Pass	
						Thiamethoxam	0.10	0.20	ND	Pass	
						Trifloxystrobin	0.10	0.20	ND	Pass	M2

Instrument: LC-QQQ

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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 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(I)(1)(b), but the sample's target analytes were not detected above the maximum allowable concentrations for the analytes in the sample.