Certificate of Analysis

Genesis Bioceuticals, LLC

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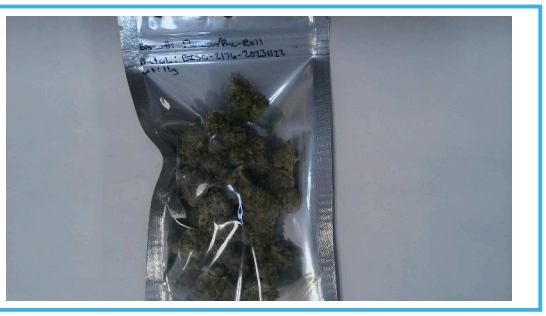
(847) 682-4899
Lic. #00000058DCQU00115543
Harvest Dates: 11/22/2023

Biscotti Flower/Pre-Roll

Plant, Bulk Flower, Extraction Method: Indoor
Dispensary License #: ; Manufacturing License #: ; Cultivation License #:

Sample: 2312TLL0069.0186

Strain: Biscotti
Parent Batch #: ; Batch#: BISC-2176-20231122; Batch Size: 11 g
Sample Received: 12/15/2023; Report Created: 12/21/2023; Expires: 12/21/2024
Manufacturing Date:
Sampling: ; Environment:



Safety

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

Cannabinoids

TPL_Potency_01

19.04%	<LOQ	NT
Total THC	Total CBD	Moisture

Analyte	LOQ	Mass	Mass	Qualifier
	%	%	mg/g	
THCa	0.20	21.19	211.9	
Δ9-THC	0.20	0.46	4.6	
Δ8-THC	0.20	ND	ND	
THCV	0.20	<LOQ	<LOQ	
CBDa	0.20	<LOQ	<LOQ	
CBD	0.20	ND	ND	
CBDV	0.20	ND	ND	
CBN	0.20	ND	ND	
CBGa	0.20	0.76	7.6	
CBG	0.20	<LOQ	<LOQ	
CBC	0.20	ND	ND	
Total		**22.41**	**224.1**	

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	Apple

Analyte	LOQ	Mass	Mass	Qualifier
	%	%	mg/g	
α-Humulene		1.1080	11.080	Q3
β-Caryophyllene		0.9090	9.090	Q3
Nerolidol		0.4800	4.800	Q3
trans-Nerolidol		0.4800	4.800	Q3
δ-Limonene		0.4630	4.630	Q3
α-Bisabolol		0.4130	4.130	Q3
Linalool		0.4040	4.040	Q3
β-Pinene		0.1760	1.760	Q3
β-Myrcene		0.1170	1.170	Q3
α-Pinene		0.1080	1.080	Q3
Terpinolene		0.0780	0.780	Q3
γ-Terpinene		0.0670	0.670	Q3
Eucalyptol		0.0480	0.480	Q3
Caryophyllene Oxide		0.0290	0.290	Q3
Camphene		0.0220	0.220	Q3
3-Carene		<	<	Q3
α-Terpinene		<	<	Q3
cis-Nerolidol		<	<	Q3
Geraniol		<	<	Q3
Guaiol		<	<	Q3
Isopulegol		<	<	Q3
Ocimene		<	<	Q3
p-Cymene		<	<	Q3
Total		**4.9020**	**49.020**	

Instrument: GCMS; Method: TPL_Terp_01
Notes:

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Brian DiMarco
Laboratory Director

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Residual Solvents — Not Tested

Analyte	LOQ	Limit	Mass	Status	Qualifier

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	ND	Pass	
Lead	500.0	1000.0	ND	Pass	
Mercury	100.0	200.0	ND	Pass	

Mycotoxins — Pass

Analyte	LOQ	Limit	Mass	Status	Qualifier
	PPB	PPB	PPB		
B1	10	20	ND	Pass	
B2	10	20	ND	Pass	
G1	10	20	ND	Pass	I1
G2	5	20	ND	Pass	
Ochratoxin A	10	20	ND	Pass	I1, R1
Total Aflatoxins	10	20	ND	Pass	

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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: ; Method:

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Pesticides TPL_Pesticides_01

Pass

Analyte	LOQ	Limit	Mass	Status	Qualifier
	PPM	PPM	PPM		
Abamectin	0.25	0.50	ND	Pass	
Acephate	0.20	0.40	ND	Pass	
Acetamiprid	0.10	0.20	ND	Pass	
Aldicarb	0.02	0.40	ND	Pass	
Azoxystrobin	0.10	0.20	ND	Pass	
Bifenazate	0.10	0.20	ND	Pass	
Bifenthrin	0.10	0.20	ND	Pass	
Boscalid	0.20	0.40	ND	Pass	M1
Carbaryl	0.10	0.20	ND	Pass	
Carbofuran	0.10	0.20	ND	Pass	
Chlorantraniliprole	0.10	0.20	ND	Pass	M1
Chlorfenapyr	0.50	1.00	ND	Pass	I1, M2
Chlorpyrifos	0.10	0.20	ND	Pass	M2
Clofentezine	0.10	0.20	ND	Pass	
Cyfluthrin	0.50	1.00	ND	Pass	
Cypermethrin	0.50	1.00	ND	Pass	
Daminozide	0.50	1.00	ND	Pass	
Diazinon	0.10	0.20	ND	Pass	
Dichlorvos	0.05	0.10	ND	Pass	I1
Dimethoate	0.10	0.20	ND	Pass	
Ethoprophos	0.10	0.20	ND	Pass	M1
Etofenprox	0.20	0.40	ND	Pass	
Etoxazole	0.10	0.20	ND	Pass	
Fenoxycarb	0.10	0.20	ND	Pass	
Fenpyroximate	0.20	0.40	ND	Pass	
Fipronil	0.20	0.40	ND	Pass	M1
Flonicamid	0.50	1.00	ND	Pass	
Fludioxonil	0.20	0.40	ND	Pass	M1

Analyte	LOQ	Limit	Mass	Status	Qualifier
	PPM	PPM	PPM		
Hexythiazox	0.50	1.00	ND	Pass	M2
Imazalil	0.10	0.20	ND	Pass	
Imidacloprid	0.20	0.40	ND	Pass	
Kresoxim Methyl	0.20	0.40	ND	Pass	
Malathion	0.10	0.20	ND	Pass	M1
Metalaxyl	0.10	0.20	ND	Pass	
Methiocarb	0.10	0.20	ND	Pass	M1
Methomyl	0.10	0.40	ND	Pass	
Myclobutanil	0.10	0.20	ND	Pass	M1
Naled	0.20	0.50	ND	Pass	
Oxamyl	0.50	1.00	ND	Pass	
Paclobutrazol	0.20	0.40	ND	Pass	
Permethrin	0.10	0.20	ND	Pass	
Phosmet	0.10	0.20	ND	Pass	M1
Piperonyl Butoxide	1.00	2.00	<LOQ	Pass	M2
Prallethrin	0.10	0.20	ND	Pass	
Propiconazole	0.20	0.40	ND	Pass	
Propoxur	0.10	0.20	ND	Pass	M1
Pyrethrins	0.42	1.00	ND	Pass	M1
Pyridaben	0.10	0.20	ND	Pass	M2
Spinosad	0.10	0.20	ND	Pass	
Spiromesifen	0.10	0.20	ND	Pass	
Spirotetramat	0.10	0.20	ND	Pass	
Spiroxamine	0.20	0.40	ND	Pass	
Tebuconazole	0.20	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-QQQ

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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