

## **Certificate of Analysis**

Company: Flavorline Cannabis 543 Belle Vista Dr Jay, VT 05859 Customer ID: 210707-1 Grower License #: SCLT0096 Sample ID: Harvest Lot Lot: SCLT0096-001 Matrix: Flower Date Sampled: N/A Date Received: 1/5/2023

Report Date: 1/13/2023 Date Analyzed: 1/12/2023 Analyst: 45 Report ID: C230105AO

## Pesticides/Mycotoxins Summary

Category II Residual Pesticide	LOQ (ppm)	Concentration (ppm)
Abamectin	0.0100	<lod< th=""></lod<>
Acephate	0.0010	<loq< th=""></loq<>
Acequinocyl	0.0010	<loq< th=""></loq<>
Azoxystrobin	0.0010	<lod< th=""></lod<>
Bifenazate	0.0010	<lod< th=""></lod<>
Bifenthrin	0.0010	<loq< th=""></loq<>
Carbaryl	0.0010	<lod< th=""></lod<>
Cypermethrin	0.0100	<loq< th=""></loq<>
Etoxazole	0.0010	<lod< th=""></lod<>
Imidacloprid	0.0010	<loq< th=""></loq<>
Myclobutanil	0.0010	<loq< th=""></loq<>
Pyrethrin I	0.0010	<lod< th=""></lod<>
Pyrethrin II	0.0010	<loq< th=""></loq<>
Spinosyn A	0.0010	<lod< th=""></lod<>
Spinosyn D	0.0010	<lod< th=""></lod<>

Category II Mycotoxin	LOQ (ppm)	Concentration (ppm)
Ochratoxin A	0.0020	NOT TESTED
Aflatoxin B1	0.0002	NOT TESTED
Alfatoxin B2	0.0010	NOT TESTED
Alfatoxin G1	0.0002	NOT TESTED
Alfatoxin G2	0.0010	NOT TESTED

Category I Residual Pesticide	LOQ (ppm)	Concentration (ppm)
Chlorpyrifos	0.0010	<loq< th=""></loq<>
Imazalil	0.0010	<loq< th=""></loq<>



12.74%
Percent Moisture

LOQ = The lowest quantity this method can reliably detect. Any pesticide or mycotoxins that was not detected is assumed to be less than the stated LOQ (<LOQ).

All results reflect dry weight of material, based on % moisture of the sample.

ppb = parts per billion

Pesticides/Mycotoxin Methodology: Liquid Chromatography with Tandem Mass Spectrometry using PerkinElme QSight® LX50 UHPLC and QSight 220 Mass Spectrometer

All moisture analysis is determined by loss-on-drying measurement using OHAUS Model MB90 Moisture Content Readers.

Certified by: \_\_\_\_

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Luke Emerson Mason (Laboratory Director, Bia Diagnostics)

This report shall not be reproduced except in full without approval of the laboratory. This is to provide assurance that parts of a report are not taken out of context. Results apply to the samples as received.

(802) 540-0148 laboratory@biadiagnostics.com