

Analytical Test Report

Client: Resinate, Inc. 120 Gilboa Street Douglas, MA 01516	Final Report MCR-S21-58554 Rev.01.00	Laboratory: MCR Labs 85 Speen St. Lower Level Framingham, MA 01701 508-872-6666
	Report Date: 29 SEPTEMBER 2021	
	METRC Tag: 1A40A01000020D2000012377 METRC Source Tag: 1A40A01000020D2000012044	

Sample ID #	Sample Name	Batch	Matrix	Date Received	Date Tested	Sample Weight
MCR-S21-58554	Critical Mass Shatter	LB9	Concentrate	24 September 2021	25-29 September 2021	4.8 g

The test results presented in this report are accurate, complete, and compliant with the MCR Labs quality control criteria.

Authorization



Andy Moy
Data Manager



This test is accredited under the laboratory's ISO/IEC 17025:2017 accreditation issued by ANSI National Accreditation Board. Refer to certificate and scope of accreditation AT-1853

Case Narrative:

This sample was received by MCR Labs from a RMD agent in a sealed container. For cannabinoids, the sample was extracted using organic solvents and analyzed via High Performance Liquid Chromatography (HPLC-UV). For microbiological contaminants, the sample was prepared using cultured enrichments, was incubated for set periods of time, and analyzed via an automated Most Probable Number (MPN) methodology. For pathogenic bacterial contaminants, the sample was analyzed via a quantitative Polymerase Chain Reaction (qPCR). Pathogenic screen includes all six STEC stains, including O157. For mycotoxin contaminants, the sample was extracted using organic solvents and analyzed via enzyme-linked immunosorbent assay (ELISA). For heavy metals, the sample was extracted using nitric acid and microwave digestion, and analyzed via Inductively Coupled Plasma Mass Spectrometry (ICP-MS). For volatile organic compounds, the sample was analyzed via Gas Chromatography – Flame Ionization Detection with Headspace Autosampler (GC-FID) using full evaporative technique. The collected data was compared to data collected from analytical reference standards at known concentrations. Unless specified by regulation, measurement uncertainty is not taken into account when reporting results and making a statement of conformity. Values reported below quantitation limits are for informational purposes.

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Requested Testing:

Test	Code	Procedure	Analytes Tested	Disposition
Cannabinoid Profile	CN	MCR-TM-0011	CBDVA, CBDV, CBDA, CBGA, CBG, CBD, THCV, THCVA, CBCV, CBN, CBNA, D9-THC, D8-THC, CBL, THCA, CBC, CBCA, CBLA, CBT	N/A
Microbiological Screen	MB	MCR-TM-0006 MCR-TM-0012	Bacterial (Total Aerobic, Total Coliform, Bile-Tolerant Gram Negative), Yeast and Mold, Pathogenic (E. coli, Salmonella)	Pass
Mycotoxin Screen	MY	MCR-TM-0015	Aflatoxin B1, Aflatoxin B2, Aflatoxin G1, Aflatoxin G2, Ochratoxin A	Pass
Heavy Metals Screen	HM	MCR-TM-0008	Arsenic (As), Cadmium (Cd), Lead (Pb), Mercury (Hg)	Pass
Volatile Organics Screen	VC	MCR-TM-0007	n-Butane	Pass

Cannabinoid Profile [MCR-TM-0011]

Analyst: JM/VB/JMW

Test Date: 26 Sep 21

The sample was analyzed for cannabinoids via High Performance Liquid Chromatography (HPLC-UV). The collected data was compared to data collected from certified analytical reference standards at known concentrations.

Table 1 - S21-58554 Critical Mass Shatter LB9 Concentrate Cannabinoid Testing

Analyte	Cannabinoid	Conc. (weight %)	Conc. (mg/g)	LOQ (weight %)	LOD (weight %)
CBDVA	Cannabidivarinic acid	ND	ND	0.10%	0.01%
CBDV	Cannabidivarin	ND	ND	0.10%	0.02%
CBDA	Cannabidiolic acid	0.1%	1.0	0.10%	0.02%
CBGA	Cannabigerolic acid	0.9%	9.0	0.10%	0.02%
CBG	Cannabigerol	0.3%	3.0	0.10%	0.04%
CBD	Cannabidiol	ND	ND	0.10%	0.03%
THCV	Tetrahydrocannabivarin	ND	ND	0.10%	0.01%
THCVA	Tetrahydrocannabivarinic acid	ND	ND	0.10%	0.03%
CBCV	Cannabichromevarin	ND	ND	0.10%	0.01%
CBN	Cannabinol	ND	ND	0.10%	0.01%
CBNA	Cannabinolic acid	0.6%	6.0	0.10%	0.01%
Δ 9-THC	Δ 9-Tetrahydrocannabinol	7.6%	76.0	0.10%	0.02%
Δ 8-THC	Δ 8-Tetrahydrocannabinol	ND	ND	0.10%	0.02%
CBL	Cannabicyclol	ND	ND	0.10%	0.02%
THCA	Tetrahydrocannabinolic acid	76.6%	766.0	0.10%	0.01%
CBC	Cannabichromene	0.1%	1.0	0.10%	0.01%
CBCA	Cannabichromenic acid	0.4%	4.0	0.50%	0.05%
CBLA	Cannabicyclolic acid	ND	ND	0.10%	0.01%
CBT	Cannabicitran	ND	ND	0.10%	0.02%

Note: There are no limits established by the Massachusetts Department of Public Health for cannabinoid concentrations. ND = Not Detected. LOQ = Limit of Quantitation. LOD = Limit of Detection.

Microbiological Screen [MCR-TM-0006] Analyst: VF Test Date: 25-28 Sep 21

The sample was analyzed for microbiological contaminants via an automated Most Probable Number (MPN) methodology with cultured enrichments.

Table 2 - S21-58554 Critical Mass Shatter LB9 Concentrate Microbiological Testing

Test ID	Test Analysis	Results	Unit	Limits	Disposition
21-58554-AC	Total Viable Aerobic Bacteria	<100	CFU/g	10 ⁴ CFU/g	Pass
21-58554-YM	Total Yeast and Mold	<100	CFU/g	10 ³ CFU/g	Pass
21-58554-CC	Total Coliforms	<100	CFU/g	10 ² CFU/g	Pass
21-58554-EB	Total Bile-Tolerant Gram Negative Bacteria	<100	CFU/g	10 ² CFU/g	Pass

Note: CFU = colony forming unit. Testing limits established by the Massachusetts Department of Public Health, Protocol for Sampling and Analysis of Finished Medical Marijuana Products and Marijuana-Infused Products for Massachusetts Registered Medical Marijuana Dispensaries, Exhibit 6.

Pathogenic Bacterial Screen [MCR-TM-0012] Analyst: MK Test Date: 26 Sep 21

The sample was analyzed for pathogenic bacterial contamination via a quantitative Polymerase Chain Reaction (qPCR).

Table 3 - S21-58554 Critical Mass Shatter LB9 Concentrate Pathogen Testing

Test ID	Test Analysis	Result	Units	Limits	Disposition
S21-58554-ECPT	STEC	Not Detected	N/A	Not Detected in 1g	Pass
S21-58554-SPT	Salmonella	Not Detected	N/A	Not Detected in 1g	Pass

Note: Testing limits established by the Massachusetts Department of Public Health, Protocol for Sampling and Analysis of Finished Medical Marijuana Products and Marijuana-Infused Products for Massachusetts Registered Medical Marijuana Dispensaries, Exhibit 6. NT = Not tested. STEC = Shiga Toxin producing E. coli

Mycotoxin Screen [MCR-TM-0015] Analyst: JG/SW Test Date: 29 Sep 21

The sample was extracted using organic solvents and analyzed via enzyme-linked immunosorbent assay (ELISA). The collected data was compared to data collected from analytical reference standard at known concentrations.

Table 4 - S21-58554 Critical Mass Shatter LB9 Concentrate Mycotoxin Testing

Test ID	Test Analysis	Result	LOD (ppb)	LOQ (ppb)	Limits (ppb)	Disposition
S21-58554-MY	Mycotoxin	Not Detected	10	10	20	Pass

Note: ND = Not Detected; LOD = Limit of Detection; LOQ = Limit of Quantitation; ppb = part per billion. Testing limits established by the Massachusetts Department of Public Health, Protocol for Sampling and Analysis of Finished Medical Marijuana Products and Marijuana-Infused Products for Massachusetts Registered Medical Marijuana Dispensaries, Exhibit 6.

Heavy Metals Screen [MCR-TM-0008] Analyst: EK/AI Test Date: 26 Sep 21

The sample was analyzed via Inductively Coupled Plasma Mass Spectrometry. The collected data was compared to data collected from certified analytical reference standards at known concentrations.

Table 5 - S21-58554 Critical Mass Shatter LB9 Concentrate Heavy Metal Testing

Test ID	Test Analysis	Result, ppb	LOD ppb	LOQ ppb	Limits ppb	Disposition	Limits (ingestion) ppb	Disposition (ingestion)
S21-58554-HM	Arsenic	ND	65.5	198.6	200	Pass	1500	Pass
S21-58554-HM	Cadmium	ND	37.6	114.0	200	Pass	500	Pass
S21-58554-HM	Mercury	ND	25.7	77.9	100	Pass	1500	Pass
S21-58554-HM	Lead	ND	40.6	123.0	500	Pass	1000	Pass

Note: ND = Not Detected; LOD = Limit of Detection; LOQ = Limit of Quantitation; BQL = Below Quantitation Limit; ppb = part per billion. Testing limits established by the Massachusetts Department of Public Health, Protocol for Sampling and Analysis of Finished Medical Marijuana Products and Marijuana-Infused Products for Massachusetts Registered Medical Marijuana Dispensaries, Exhibit 4.

VC Screen [MCR-TM-0007]*Analyst: IM**Test Date: 26 Sep 21*

The sample was analyzed via Gas Chromatography – Flame Ionization Detection with Headspace Autosampler. The collected data was compared to data collected from certified analytical reference standards at known concentrations.

Table 6 - S21-58554 Critical Mass Shatter LB9 Concentrate Residual Solvent Testing

Test ID	Analyte	Result (ppm)	LOD (ppm)	LOQ (ppm)	Limits (ppm)	*USP Result (ppm)	Disposition
S21-58554-VC	n-Butane	ND	0.29	5	12	ND	Pass

Note: ND = Not Detected; LOD = Limit of Detection; LOQ = Limit of Quantitation; BQL = Below Quantitation Limit; ppm = part per million. Testing limits established by the Massachusetts Department of Public Health, Protocol for Sampling and Analysis of Finished Medical Marijuana Products and Marijuana-Infused Products for Massachusetts Registered Medical Marijuana Dispensaries, Exhibit 7. The uncertainty budget for propane is 0.12 ppm; n-Butane - 0.10 ppm; Ethanol - 0.15 ppm. *USP 34 General Notices 7.20.

QA/QC

Cannabinoid Profile [MCR-TM-0011]

Analyst: YD/KM

Test Date: 26 Sep 21

The sample data for certified reference standards was collected at known concentrations of cannabinoids in solution.

QC-0.025 mg/mL 19 cannabinoid multi-component 9/7/2021

ID	Cannabinoid	Nominal Prep Conc (mg/mL)	Measured Conc. (mg/mL)	Recovery (%)
CBDVA	Cannabidivarinic acid	0.025	0.026	104%
CBDV	Cannabidivarin	0.025	0.025	100%
CBDA	Cannabidiolic acid	0.025	0.025	100%
CBGA	Cannabigerolic acid	0.025	0.025	100%
CBG	Cannabigerol	0.025	0.024	96%
CBD	Cannabidiol	0.025	0.025	100%
THCV	Tetrahydrocannabivarin	0.025	0.025	100%
THCVA	Tetrahydrocannabivarinic acid	0.025	0.025	100%
CBCV	Cannabichromevarin	0.025	0.025	100%
CBN	Cannabinol	0.025	0.026	104%
CBNA	Cannabinolic acid	0.025	0.026	102%
Δ9-THC	Δ9-Tetrahydrocannabinol	0.025	0.026	104%
Δ8-THC	Δ8-Tetrahydrocannabinol	0.025	0.026	104%
CBL	Cannabicyclol	0.025	0.026	104%
THCA	Tetrahydrocannabinolic acid	0.025	0.026	104%
CBC	Cannabichromene	0.025	0.025	100%
CBCA	Cannabichromenic acid	0.025	0.026	104%
CBLA	Cannabicyclic acid	0.025	0.025	100%
CBT	Cannabicitran	0.025	0.025	100%

Criteria for successful analysis is QC recovery to be ≤20% above or below nominal.

Microbiological Screen [MCR-TM-0006]

Analyst: TM/DGM/IM

Test Date: 08 Sep 21

Quality control checks are performed to confirm that the equipment used for reading incubated microbiological cultures, which are done at various concentrations, are working correctly and that the fluorescence readings are accurate. QC checks are performed within 30 days of the recorded measurements.

Date of most recent QC check: Tempo2 QC 09/8/2021
 Status: Pass

Pathogenic Bacterial Screen [MCR-TM-0012]

Analyst: MK

Test Date: 26 Sep 21

Quality control checks are performed to validate the equipment used for reading incubated pathogenic bacterial cultures. QC checks are run with every analysis.

Date	QC Check	Pathogen	Result	Disposition
9/26/2021	Control (+)	STEC	Positive	Pass
9/26/2021	Control (-)	STEC	Negative	Pass
9/26/2021	Control (+)	Salmonella	Positive	Pass
9/26/2021	Control (-)	Salmonella	Negative	Pass

Mycotoxin Screen [MCR-TM-0015]

Analyst: JG/SW

Test Date: 29 Sep 21

Solutions were spiked with toxin reference materials at given concentrations and tested for toxin presence.

QC Sample	Result	Disposition
Negative Control	Negative	Pass
Positive Control	Positive	Pass

Heavy Metals Screen [MCR-TM-0008]

Analyst: EK/AI

Test Date: 26 Sep 21

QC samples were prepared at target concentrations and injected at the end of the sequence.

Analyte	Prepared analyte concentration, ppb	Analyte measured, ppb	QC recovery (%)
Arsenic (As)	1.00	0.94	94%
Cadmium (Cd)	1.00	0.97	97%
Mercury (Hg)	0.50	0.44	88%
Lead (Pb)	3.00	2.72	91%

Criteria for successful analysis is QC recovery to be $\leq 20\%$ above or below nominal.

VC Screen [MCR-TM-0007]

Analyst: IM

Test Date: 26 Sep 21

A QC sample was prepared at a known concentration and injected.

Analyte	µg analyte detected	Nominal analyte, µg	Recovery
Hexane	4.5	5	90%

Criteria for successful analysis is QC recovery to be ≤30% above or below nominal.

END OF REPORT