

CBD Soap

Lab ID: 1910250-02

BBPS LLC
AG-R1050457IHH

METRC Batch ID:

Date Sampled: 10/31/19

Date Printed: 11/1/19

Report cannot be used for OLCC/OHA compliance.

Potency Analysis

Analytical Method: De Backer, Journal of Chromatography b.2009. 11.004 - SOP 102

Cannabinoids	mg/g	LOQ
THCA	0.0881	0.0148
delta 9-THC	0.0356	0.0148
delta 8-THC	0.0289	0.0148
CBGA	< LOQ	0.0148
CBDA	0.366	0.0148
CBD	0.342	0.0148
CBN	0.0210	0.0148
CBG	< LOQ	0.0148
CBC	0.0891	0.0148

Total THC
0.0356 mg/g

Total CBD
0.342 mg/g

<LOQ - Results below the Limit of Quantitation

Acid form of THC/CBD are decarboxylated by heat, lose 12% of original mass as CO2. Result = *bioactive*

"Total" Cannabinoid accounts for decarboxylation and moisture content. Total THC = [(THCA*0.877) + Δ9THC] / (100%-MC)



Erik Werstler
Lab Director

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Quality Control Potency

Batch: B19J174 - Potency

Blank(B19J174-BLK1)

Analyte	Result	LOQ	Units	%Recovery Limits	Extracted	Analyzed	Notes
THCA	< LOQ	0.150	mg/g		10/31/19 15:46	10/31/19 18:29	
delta 9-THC	< LOQ	0.150	mg/g		10/31/19 15:46	10/31/19 18:29	
CBGA	< LOQ	0.150	mg/g		10/31/19 15:46	10/31/19 18:29	
CBDA	< LOQ	0.150	mg/g		10/31/19 15:46	10/31/19 18:29	
CBD	< LOQ	0.150	mg/g		10/31/19 15:46	10/31/19 18:29	
CBN	< LOQ	0.150	mg/g		10/31/19 15:46	10/31/19 18:29	
CBG	< LOQ	0.150	mg/g		10/31/19 15:46	10/31/19 18:29	
delta 8-THC	< LOQ	0.150	mg/g		10/31/19 15:46	10/31/19 18:29	
CBC	< LOQ	0.150	mg/g		10/31/19 15:46	10/31/19 18:29	

LCS(B19J174-BS1)

Analyte	% Recovery	LOQ	Units	%Recovery Limits	Extracted	Analyzed	Notes
THCA	92.5	0.146	mg/g	85-115	10/31/19 15:46	10/31/19 18:48	
delta 9-THC	102	0.146	mg/g	85-115	10/31/19 15:46	10/31/19 18:48	
CBDA	106	0.146	mg/g	85-115	10/31/19 15:46	10/31/19 18:48	
CBD	108	0.146	mg/g	85-115	10/31/19 15:46	10/31/19 18:48	

Notes and Definitions

- B Analyte detected in method blank, but not associated samples.
 - B2 Analyte detected in sample and associate method blank.
 - C Interference due to co-elution.
 - D Initial result exceeded calibration range, reported data are based on analysis of a dilution.
 - H Non-homogenous sample matrix affecting RPD and/or QC.
 - I Manual Integration was performed.
 - L Duplicate sample relative percent difference (RPD) exceeds QC limits.
 - M Anomalous results due to matrix interference
 - P Peaks manually split.
 - Q1 QC out of limits but still ok
 - Q2 Quality Control outside QC limits. Data considered estimate.
 - Q3 CCV was above the acceptance criteria. Non-detect samples are considered acceptable.
 - Q4 CCV was below the acceptance criteria, however the sample still exceeds the regulatory limit.
 - R Marginal Exceedence.
 - U Reported result is an estimate. The analyte was detected above the calibration range.
 - X Problems with initial analysis, reported data are from reinjection of prepared sample.
- <LOQ - Results below the Limit of Quantitation - Compound not detected



Erik Werstler
Lab Director