

Certificate of Analysis

Kaycha Labs

N/A Matrix: Edible

Sample:M000409002-001 Harvest/Lot ID: 02 Seed to Sale #N/A Batch Date :N/A Batch#: 0029 Sample Size Received: 3.5 gram Retail Product Size: 3.5 Ordered : 04/07/20 Sampled : 04/07/20 Completed: 04/10/20 Expires: 04/10/21 Sampling Method: SOP Client Method

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Apr 10, 2020 | Reserve Infusibles

South Carolina, United States 29801

PRODUCT IMAGE SAFETY RESULTS



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Total THC



NOT



NOT TESTED



cotoxins Residuals



ilth Water Activity

NOT TESTED

y Moisture

NOT TESTED



NOT TESTED

MISC.

NOT TESTED

CANNABINOID RESULTS



0.000% THC/Container :0.000 mg



Solvents

NOT TESTED



D9-THC THCA CBD CBDA D8-THC THCV CBN CBDV CBC CBG CBGA ND ND 0.270% ND ND ND ND ND ND ND ND 2.700 ND mg/g 0.01 0.01 0.01 0.01 0.01 0.01 0.01 0.01 0.01 0.01 0.01 LOD % % % % % % % % %

Cannabinoid Profile Test

Analyzed by	Weight	Extraction date :	Extracted By :
1	1g	NA	NA

 Analysis Method -SOP.T.40.020, SOP.T.30.050
 Reviewed On - 04/10/20 08:59:35

 Analytical Batch -M0000434POT
 Instrument Used : HPLC Potency Analyzer Batch Date : 04/09/20 14:09:24

Reagent Dilution Consums. ID

Full spectrum cannabinoid analysis utilizing High Performance Liquid Chromatography with UV detection (HPLC-UV). (Method: SOP.T.30.050 for sample prep and Shimadzu High Sensitivity Method SOP.T.40.020 for analysis. LOQ for all cannabinoids is 1 mg/L). Measurement of Uncertainty: 2.7%

This report shall not be reproduced, unless in its entirety, without written approval from Kaycha Labs. This report is an Kaycha Labs certification. The results relate only to the material or product analyzed. Test results are confidential unless explicitly waived otherwise. Void after 1 year from test end date. Cannabinoid content of batch material may vary depending on sampling error. IC=In-control QC parameter, NC=Non-controlled QC parameter, ND=Not Detected, NA=Not Analyzed, ppm=Parts Per Million, pb=Parts Per Billion. Limit of Detection (LoD) and Limit Of Quantitation (LoQ) are terms used to describe the smallest concentration that can be reliably measured by an analytical procedure. RPD=Reproducibility of two measurements. Action Levels are State determined thresholds for human safety for consumption and/or inhalation. The result >99% are variable based on uncertainty of measurement (UM) for the analyte. The UM error is available from the lab upon request. The "Decision Rule" for the pass/fail does not include the UM. The limits are based on F.S. Rule 64-4.310.

David Greene

Lab Director State License # 19-05-02P ISO Accreditation # 17025:2017 Dendon

04/10/2020

Signature

Signed On