



# Certificate of Analysis

Sample:KN20822009-001  
Harvest/Lot ID: CWE-SG-100318-2  
Batch#: CWE-SG-100318-2  
Seed to Sale# N/A  
Batch Date: 10/03/18  
Sample Size Received: 5 units  
Total Batch Size: N/A  
Retail Product Size: 5 units  
Ordered : 08/17/22  
Sampled : 08/17/22  
Completed: 08/25/22  
Sampling Method: N/A

**PASSED**

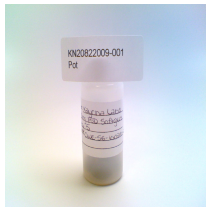
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Aug 25, 2022 | Commonwealth Extracts, LLC

6900 Riverport Dr  
Louisville, KY, 40258, US



PRODUCT IMAGE



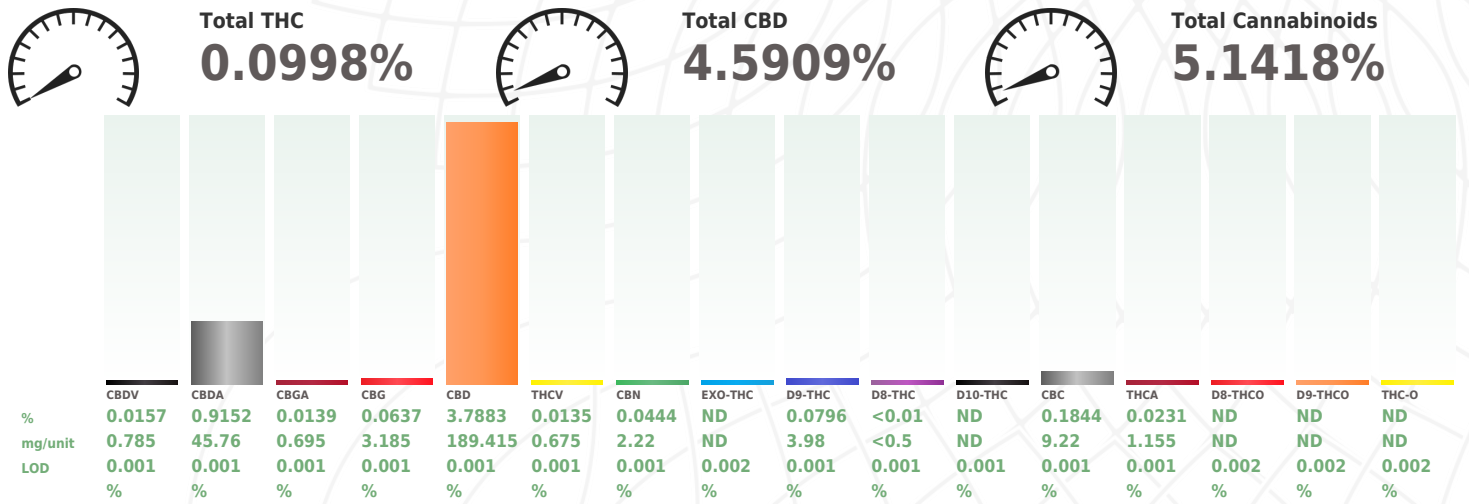
per unit

SAFETY RESULTS

 Pesticides NOT TESTED	 Heavy Metals NOT TESTED	 Microbials NOT TESTED	 Mycotoxins NOT TESTED	 Residuals Solvents NOT TESTED	 Filth NOT TESTED	 Water Activity NOT TESTED	 Moisture NOT TESTED	 Terpenes NOT TESTED
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MISC.

 **Cannabinoid** **PASSED**



Analyzed by: 2692      Weight: 0.2130g      Extraction date: 08/22/22 15:20:10      Extracted by: 2692

Analysis Method : Expanded Measurement of Uncertainty: Flower Matrix d9-THC:12.7%, THCA: 9.5%, TOTAL THC 11. 1%. These uncertainties represent an expanded uncertainty expressed at approximately the 95% confidence level using a coverage factor k=2 for a normal distribution.

Analytical Batch : KN002806POT      Reviewed On : 08/24/22 15:34:38  
Instrument Used : HPLC E-SHI-008      Batch Date : 08/22/22 09:20:06  
Running on : N/A

Dilution : N/A  
Reagent : 062422.02; 063022.R01; 063022.R02  
Consumables : 294033242; n/a; 12265-115CC-115  
Pipette : E-GIL-010; E-EPP-081

Full spectrum cannabinoid analysis utilizing High Performance Liquid Chromatography with UV/PDA detection (HPLC-UV/PDA). (Method: SOP.T.30.031.TN for sample prep and Shimadzu High Sensitivity Method SOP.T.40.020 for analysis). \*Based on FL action limits.

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, ppb=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.

**Sue Ferguson**

Lab Director

State License # n/a  
ISO Accreditation # 17025:2017

  
Signature

08/25/22

Signed On