



Certificate of Analysis

Sample:KN20225006-003

Harvest/Lot ID: 2201

Batch#: 5010-2201

Seed to Sale# N/A

Batch Date: 02/03/22

Sample Size Received: 4g gram

Total Weight/Volume: N/A

Retail Product Size: .85 gram

ordered : 02/18/22

sampled : 02/18/22

Completed: 03/01/22 Expires: 03/01/23

Sampling Method: SOP Client Method

PASSED

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Mar 01, 2022 | The TabEASE Company

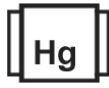
1005 Perkins Ave
Waukesha, WI, 53186-5249, US



PRODUCT IMAGE SAFETY RESULTS



Pesticides
NOT TESTED



Heavy Metals
NOT TESTED



Microbials
NOT TESTED



Mycotoxins
NOT TESTED



Residuals Solvents
NOT TESTED



Filtration
NOT TESTED



Water Activity
NOT TESTED



Moisture
NOT TESTED



Terpenes
NOT TESTED

MISC.

CANNABINOID RESULTS



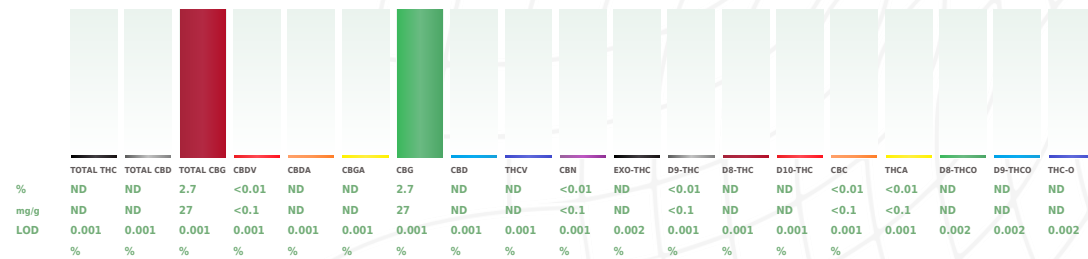
Total THC
ND



Total CBG
2.70%



Total Cannabinoids
2.70%



Cannabinoid Profile Test

Analyzed by: 113 Weight: 0.2052g Extraction date: 02/25/22 05:02:32 Extracted By: 113
 Analysis Method -Expanded Measurement of Uncertainty: Flower Matrix d9-THC:12.7%, THCa: 9.5%, TOTAL THC 11.3%. These uncertainties represent an expanded uncertainty expressed at approximately the 95% confidence level using a coverage factor k=2 for a normal distribution. Reviewed On - 03/01/22 10:46:06
 Analytical Batch -KN002015POT Instrument Used : HPLC E-SHI-008 Running On : Batch Date : 02/25/22 12:13:36

Reagent	Dilution	Consumables ID
081321.R04 022522.R01 021622.R03	40	947.251 12123-046CC-046

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.40.031 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

Sue Ferguson
Signature

03/01/22

Signed On