



Certificate of Analysis



Honey Extra Strength

Matrix: Edible

Accession Number: 051021UD0017

Harvest/Lot ID:

Seed to Sale: *

Batch Date: 04/21/21

Batch #:

Sample Size Received: 1 units

Retail Product Size: 180 units

Ordered: 04/21/21

Completed: 05/13/21

Expires: 05/12/22

Sampling Method: SOP Client Method

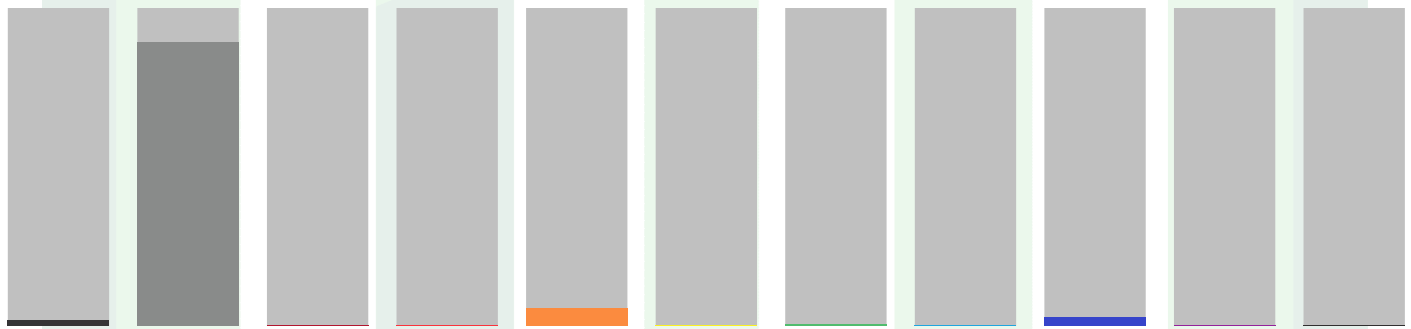
May 13, 2021 | Natural Organix
dba Beezy Beez



Brooklyn, NY,
(718) 676-9454

CANNABINOID RESULTS

Total THC 0.019% THC/Container :34.2 mg	Total CBD 0.658% CBD/Container :1184.4 mg	Total Cannabinoids 0.736% Cannabinoids/Container :1324.8 mg
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Cannabinoid	Percentage	mg/g	LOD
CBC	0.013%	0.130 mg/g	0.001
CBD	0.658%	6.580 mg/g	0.0001
CBDA	ND	ND	0.001
CBDV	0.002%	0.020 mg/g	0.001
CBG	0.041%	0.410 mg/g	0.001
CBGA	ND	ND	0.001
CBN	0.003%	0.030 mg/g	0.001
D8-THC	ND	ND	0.001
D9-THC	0.019%	0.190 mg/g	0.0001
THCA	ND	ND	0.001
THCv	ND	ND	0.001

Full spectrum cannabinoid analysis utilizing High Performance Liquid Chromatography with UV detection (HPLC-PDA). (Method: SOP.KY.02.005) sample prep and Shimadzu High Sensitivity Method SOP.KY.02.012 for analysis. LOQ for all cannabinoids is 1 mg/L. % = %w/w = Percent (Weight of Analyte/Weight Product) Total Cannabinoids result reflects the absolute sum of all cannabinoids detected. **Total Potential THC/CBD is calculated using the following formulas to take into account the loss of a carboxyl group during decarboxylation Total THC = THC + (THCa*0.877) Total CBD = CBD + (CBDa*0.877) null

This report shall not be reproduced, unless in its entirety, without written approval from Universal Diagnostics. This report is an Universal Diagnostics 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.

David Greene
Lab Director
State License # 19-05-02P
ISO Accreditation # PJLA
ISO17025

Signature

05/13/21
Signed On