



# Certificate of Analysis



chocolate piece  
Matrix: Edible  
Accession Number: 032221UD0005  
Harvest/Lot ID: Delta 8 Batch  
Seed to Sale: \*  
Batch Date: 03/17/21  
Batch #: 316-215  
Sample Size Received: 5 units  
Retail Product Size: 1 units  
Ordered: 03/17/21  
Completed: 03/27/21  
Expires: 03/26/22  
Sampling Method: SOP Client Method

Mar 27, 2021 | Incentive  
Gourmet

70 Clinton Rd  
Fairfield, New Jersey, 07004  
9738828850



## CANNABINOID RESULTS

<b>Total THC</b> <b>0.004%</b>	<b>Total CBD</b> <b>0.000%</b>	<b>Total Cannabinoids</b> <b>0.237%</b>
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	CBC	CBD	CBDA	CBDV	CBG	CBGA	CBN	D8-THC	D9-THC	THCA	THCV
	ND	ND	ND	ND	ND	ND	ND	0.233%	0.004%	ND	ND
	ND	ND	ND	ND	ND	ND	ND	2.330 mg/g	0.040 mg/g	ND	ND
<b>LOD</b>	<b>0.001</b>	<b>0.0001</b>	<b>0.001</b>	<b>0.001</b>	<b>0.001</b>	<b>0.001</b>	<b>0.001</b>	<b>0.001</b>	<b>0.0001</b>	<b>0.001</b>	<b>0.001</b>

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)

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

Signature

03/27/21

Signed On