

## Certificate of Analysis

Jan 26, 2021 | IC FORMULATIONS

20221 NE 16th ST MIAMI, FL, 33179, US

IC FORMULATIONS

## Kaycha Labs

D8 Softgel N/A Matrix: Edible

Sample:DA10122009-001 Harvest/Lot ID: 1014-1 / 01.2023

> Seed to Sale #N/A Batch Date :01/15/21

Batch#: 011521 Sample Size Received: 13 units Retail Product Size: 0.55895

Ordered : 01/20/21

Sampled: 01/20/21 Completed: 01/26/21 Expires: 01/26/22 Sampling Method: SOP Client Method

**PASSED** 

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PRODUCT IMAGE

SAFETY RESULTS







Heavy Metals



Microbials



Mycotoxins



Solvents



Filth



Water Activity



Moisture

**NOT TESTED** 

Pesticides NOT TESTE

**CANNABINOID RESULTS** 

TOTAL D8 THC
2.967%
D8 THC/Softgel :16.584 m



TOTAL CBD

0.008%

TOTAL CBD/Softgel :0.045 mg

Total C 3.1 Total Ca :17.355

Total Cannabinoids
3.105%
Total Cannabinoids/Softgel:17.355 mg



## **Cannabinoid Profile Test**

Analyzed by	Weight	Extraction date :	Extracted By:
450	1.6761g	01/22/21 08:01:28	574
Analysis Method -SOP.T.40.020, SOP.T.30.050		Reviewed On - 01/25/21 12:29:59	Batch Date: 01/22/21 09:39:48
Analytical Batch -DA021476POT		Instrument Used : DA-LC-003	

 Reagent
 Dilution
 Consums. ID

 110520.67
 400
 280670723

 012021.R16
 11989-024CC-024

 012021.R15
 76262-590

 070820.25
 914C4-9144K

 929C6-0293H

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).

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 (LoO) 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.

**Jorge Segredo** 

Lab Director

State License # CMTL-0002 ISO Accreditation # ISO/IEC 17025:2017 Accreditation PJLA-Testing 97164



01/26/2021

Signature Signed On