



Certificate of Analysis




Summary COA (scan QR code for complete Certificate of Analysis)

Sample: LA40223004-008
Laboratory License # 69204305475717257553
Sample Size Received: 1 units
Retail Product Size: 30 gram
Ordered: 02/23/24
Sampled: 02/23/24
Completed: 02/28/24

Feb 28, 2024 | PX International LLC

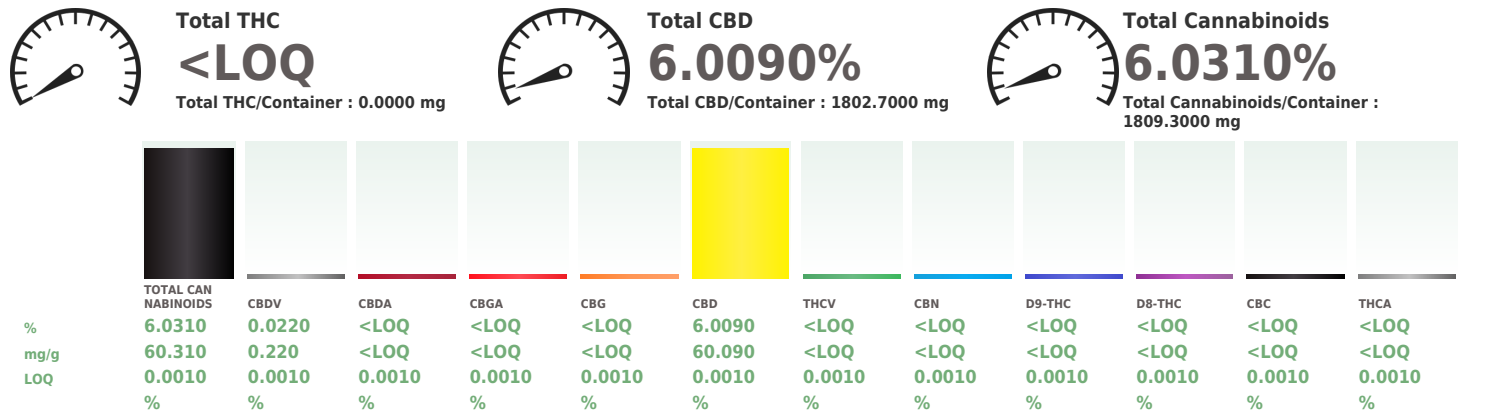
PASSED

Pages 1 of 1

PRODUCT IMAGE	SAFETY RESULTS									MISC.
										
	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	Homogeneity Testing NOT TESTED	Terpenes NOT TESTED

1 unit = 1 tincture bottle; 30g

 **Cannabinoid** **PASSED**



Analyzed by: 877, 1526 Weight: 2.8635g Extraction date: 02/28/24 09:04:55 Extracted by: 1525, 877

Analysis Method : SOP.T.30.031.NV; SOP.T.40.031.NV
Analytical Batch : LA004734POT Reviewed On : 02/28/24 11:41:10
Instrument Used : LV-SHIM-001 Batch Date : 02/26/24 14:14:56
Analyzed Date : N/A

Dilution : 400
Reagent : 120723.33; 022024.02; 021324.08; 112823.33; 020224.R09; 022124.R09
Consumables : 042c6; 258638; 265084
Pipette : LV-PIP-015; LV-PIP-008; LV-PIP-023

Cannabinoid analysis utilizing Ultra High Performance Liquid Chromatography with UV Detection (UHPLC-UV). Method SOP.T.30.031.NV for sample preparation and SOP.T.40.031.NV for analysis. Total THC = d8-THC + d9-THC + 0.877 * THCA, Total CBD = CBD + 0.877 * CBDA

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 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 NV regulations.

Kelly Zaugg
Lab Director

State License # L003
ISO 17025 Accreditation # ISO/IEC
17025:2017: 97164



Signature
02/28/24