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## CERTIFICATE OF ANALYSIS

Product Name:	UCG Rozzie Stick Strawberry Guava	Date of Issue:	17AU23
Product Code:	FPUCGVAP001	GTIN-12:	628693908009
COA Number:	COA-FPUCGVAP001- PKRS230804-01	Lot Number:	PKRS230804
Format:	1 g disposable vape pen	Packaging Date: 2023-08-04	
Equivalency:	4.0 g dried flower / unit	Expiry:	Not Established

ANALYSIS	METHOD	SPECIFICATIONS/	Results _	A CONTRACTOR OF THE PARTY OF TH	sults orm ?
Parameters		ACCEPTANCE	A-VANAGE -	YES	NO
Description	Organoleptic	Vape pen filled with a honey- like amber translucent viscous cannabis extract/liquid	Conforms	$\boxtimes$	
Foreign Matter	Magnified Visual (Product specific house method)	NMT 3%	Conforms	$\boxtimes$	
Fill Weight Variation	USP <905>	1 g net (RSD <7.0%)	Conforms		
Cannabinoid Profile*	。 《古代·阿拉尔·西斯·西斯·斯尔·斯尔·斯尔·斯尔·斯尔·斯尔·斯尔·斯尔·斯尔·斯尔·斯尔·斯尔·斯尔	(1000年) (1000年) (1000年) (1000年) (1000年)	(Kontal) place (desirati)	185 et)	nego d
Tetrahydrocannabinol (d9-THC)	HPLC-UV Validated Assay	Report % or mg/g	512.6 mg/g	$\boxtimes$	
Tetrahydrocannabinol Acid A (THCA)	HPLC-UV Validated Assay	Report % or mg/g	149.2 mg/g		
Cannabidiol (CBD)	HPLC-UV Validated Assay	Report % or mg/g	<0.1 mg/g		
Cannabidiolic Acid (CBDA)	HPLC-UV Validated Assay	Report % or mg/g	2.8 mg/g	$\boxtimes$	
Cannabigerol (CBG)	HPLC-UV Validated Assay	Report % or mg/g	46.7 mg/g		
Cannabigerolic Acid (CBGA)	HPLC-UV Validated Assay	Report % or mg/g	78.0 mg/g		
Cannabichromene (CBC)	HPLC-UV Validated Assay	Report % or mg/g	6.9 mg/g	$\boxtimes$	
Cannabidivarin (CBDV)	HPLC-UV Validated Assay	Report % or mg/g	< 1.0 mg/g		
Cannabinol (CBN)	HPLC-UV Validated Assay	Report % or mg/g	1.1 mg/g	$\boxtimes$	
Δ8-Tetrahydrocannabinol (d8-THC)	HPLC-UV Validated Assay	Report % or mg/g	1.1 mg/g		
Total THC	d9-THC % + (0.877 x THCA %)	800-860 mg/g	643.5 mg/g	$\boxtimes$	
Total CBD	CBD % + (0.877 x CBDA %)	0-10 mg/g	2.5 mg/g		



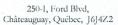
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Microbia!*	1、一层、产品的人人的政治				
Total Aerobic Microbial Count	USP and EP Harmonized Method **AOAC	100 CFU/g	< 10 CFU/g		) <u> </u>
Total Combined Yeast and Molds Count	USP and EP Harmonized Method **AOAC	10 CFU/g	< 10 CFU/g		
Bile-tolerant Gram-Negative Bacteria	USP and EP Harmonized Method **AOAC	Absent in 1g	Absent	$\boxtimes$	
Escherichia coli	USP and EP Harmonized Method **AOAC	Absent in 1g	Not required**	$\boxtimes$	
Salmonella	USP and EP Harmonized Method **AOAC	Absent in 25g	Not required**	$\boxtimes$	
Pseudomonas aeruginosa	USP and EP Harmonized Method **EP	Absent in 1g	Absent	$\boxtimes$	
Staphylococcus aureus	USP and EP Harmonized Method **EP	Absent in 1g	Absent	$\boxtimes$	
Heavy Metals*	The second of the second of the second	CHARLES CALLED A STATE OF THE S	N. FALL WOLLSAN	· pianta	is to be a
Arsenic (inorganic)	ICP-MS Validated Assay **USP / ICH	0.2 ppm	< 0.10 ppm	$\boxtimes$	
Cadmium	ICP-MS Validated Assay **USP / ICH	0.2 ppm	< 0.10 ppm		
Lead	ICP-MS Validated Assay **USP / ICH	0.5 ppm	< 0.25 ppm		
Mercury (inorganic)	ICP-MS Validated Assay **USP / ICH	0.1 ppm	< 0.05 ppm	$\boxtimes$	
Pesticides*			W. K.S. Transition of the State of		
Pesticide Residue  Mycotoxins*	GC-MS / LC-MS Validated Assay **ORG-M-051 / AOAC.2007.01	Conforms to Health Canada Pesticide List and Tolerance Limits, adapted for the finished product.	Conforms	$\boxtimes$	
Miyootoxiiio	[1] (1) (1) (1) (1) (1) (1) (1) (1) (1) (1)	· 高麗建一起的一点人员和克里克尔。这个人的	<b>建業を確認する。 ・                                    </b>	。这样单," / / / / / / / / / / / / / / / / / / /	2011年18日
	LC-MS-MS Validated Assay	EP			
Aflatoxins	**EP 2.8.18/Quantitation of	Aflatoxin B1 2 ppb	< 2.00 ppb	$\boxtimes$	
	Aflatoxins in Cannabis	Total Aflatoxins (B1, B2, G1, G2) 4 ppb	< 2.00 ppb		
Residual Solvents	2011年2月日本 日本 日本 1941年1月1日本	不是 为国际的现代,然后的特殊的 (1995年) 2000年	4. 中国的1995年1757年45日	AND A	
Residual Solvents	GC-MS Validated Assay **USP 467/Residual Solvents in Plant Extracts	5000 ppm Health Canada Tolerance Limit	<2400 ppm		

Reference: \* COA 2023-02130 Rev1, \*\* As per USP <1111> MICROBIOLOGICAL EXAMINATION OF NONSTERILE PRODUCTS: ACCEPTANCE CRITERIA FOR PHARMACEUTICAL PREPARATIONS AND SUBSTANCES FOR PHARMACEUTICAL USE. Table 1.





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Acceptance Criteria for Microbiological Quality of Nonsterile Dosage Forms - Inhalation use (special requirements apply to liquid preparations for nebulization).

I hereby attest that the above information is authentic and accurate as per third party analytical testing results. I further attest that the batch processing, packaging and analysis records were reviewed and found to be in compliance with all applicable regulations including part 5 and 6 of the Cannabis Regulations (SOR/2018-144). An adequate retention sample of this lot has been kept to support verification of quality (if necessary). This lot has not been subjected to any radiation treatment. All input materials have been evaluated and any pesticides used are in compliance with the product specifications and in accordance to the Health Canada Pesticide List and Tolerance Limits. This lot has been manufactured as per the Master Production Documents in accordance with Good Manufacturing Practices. There were no deviations or non-conformities reported for this lot release. We therefore deem this product acceptable for sale.

Marc Bergeron - QAP

QA APPROVED

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Date