

## 1st Quarterly Report - 2017

# Cannabis Flower Sampling Recommendations Derived From Analysis Of Cannabinoid Variance in a 4 Cultivar Cultivation Test Plot

This report is generated for the General and Sponsoring Members of The Clinical Endocannabinoid System Consortium (The CESC)

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# **Executive Summary**

We continue our ongoing Cannabis Chemotyping analyses with a detailed description of Cannabinoid content from a 2016 season Humboldt County test plot. The Tetrahydrocannabinolic acid (THCA) % potency results from a Design of Experiment (DOE) screening model of various cultivation main effects is presented. A sampling scheme utilizing a Sentinel Plant is proposed. Power analysis is applied to the dataset to derive optimal sampling number and guidelines for potency analysis, potentially applicable to much larger cultivation plots. Our intent is to provide some empirical background and guidance to support best practices for sampling and testing the Cannabis agricultural commodity.



# **Table of Contents**

L	st Quarterly Report – 2017	1
	Executive Summary	1
	List of Figures	2
	Background & Rationale	4
	The Cannabis industry needs robust sampling guidance	4
	Details of the Method & Approach	4
	Ad-hoc Design of Experiments (DOE); targeting main effects only	4
	Testing the Sentinel Plant Approach	6
	Strict, rigorous clonal derivation of nursery stock used in study	7
	Certain strains were not ripe at harvest	7
	Entire untrimmed, cured flower (bud) was analyzed for Cannabinoids by HPLC-DAD	8
	Key Points So Far (I):	<u>9</u>
	Study Results	10
	Can CBGA % Be Used as a Signal for When to Harvest?	10
	No evidence that strains exhibit different variances for THCA %	11
	Sampling across a Sentinel Plant should be representative of sampling across the entire grow plo	ot11
	Flower position can be a significant factor affecting THCA % (in some strains / conditions)	13
	Greenhouse conditions minimize THCA % variance contrasted with outdoor	15
	Trimming flowers can marginally increase THCA% potency	17
	Key Points So Far (II):	17
	Using Power Analysis to Determine Optimal Sampling Number	18
	Summary, Conclusions, & Recommendations	19



### **Background & Rationale**

### The Cannabis industry needs robust sampling guidance

How many of us have gone into a medical Cannabis dispensary and been told by the staff that a particular strain has, for example, 24.34% THC? What does that mean? There are typically no ranges reported for these values, no error factor, or no ± values for guidance. Does that mean that every flower in the jar has the same potency? It stretches credibility that this is the case. We intuitively

expect there is likely some range, but we really are not given any guidance.. In other industries sampling, reporting, and testing guidelines are a routine part of best practices. Because the medical Cannabis industry is just now coming out of the shadows of prohibition we lack such clear guidance. The CESC believes that the creation of these best practices should be an integral part of the medical Cannabis industry. As part of our mission, we recently undertook a study of potency variance in controlled Cannabis cultivation test plots in order to come up with recommendations for sampling for potency testing. The results we obtained and the conclusions we have reached are described in this report. It is our intention and belief that adaption of these recommendations

"Because the medical cannabis industry is just now coming out of the shadows of prohibition, we lack such clear guidance."



-Dr. John Abrams

may lead to a safer and more reliable medical Cannabis product.

The basis of our sampling study derives from a larger effort mounted by <u>SICPA</u>. This organization is a leading global provider of secured identification, traceability and authentication solutions and services. It produces the authenticity stamps on cigarette packages and is currently being evaluated for a similar role in the State of California's Medical Cannabis Track and Trace program. During the 2016 growing season, qualified cultivators took part in test plot program in conjunction with the Humboldt County Agricultural Department for MCRSA compliance. The program was designed to provide yield data from defined canopy sizes. This pilot study offered us a perfect opportunity to merge in our own sampling study. Our approach was to incorporate an *ad-hoc* Design of Experiments (DOE) Approach and analyze potency variance to derive sampling size guidelines. By using a statistical tool known as Power Analysis, we derived the number of (replicate) samples necessary for analysis. This method is based on the variance or experimental error of the data.



# **List of Figures**

Figure 1 Summary of factors studied in Cannabis test plots	5
Figure 2 Experimental Design for 4 Cultivar test plot potency determination	6
Figure 3 Plant and flower assignments within the plot.	7
Figure 4 Outline of propagation, harvest, and curing conditions used in this study	8
Figure 5 "Candyland", "Ogre", and "Tangie" Cannabis strains grown in the outdoor test plot	8
Figure 6 Outline of analysis conditions used in this study	9
Figure 7 Cannabinoid levels quantified in the test plot samples	10
Figure 8 No evidence that strains exhibit different variances for THCA %	11
Figure 9 No Evidence for Within Plant vs Between Plant Variance difference for THCA %	12
Figure 10 THCA %: Screening Main Effects Using Profile Plots	13
Figure 11 THCA %: Bud Source Depth for "Candyland" strain	14
Figure 12 THCA %: Bud Source Depth for "Tamgie" strain	14
Figure 13 THCA %: Outdoor vs. Greenhouse for "OG" Strain	15
Figure 14 "OG" Strain: Outdoor vs. Greenhouse	16
Figure 15 THCA %: Planting Position relative difference for "OG" Strain grown outdoors	16
Figure 16 THCA %: Effect of Trimming for "OG" Strain	17
Figure 17 Using Power Analysis to determine optimal sampling number	18
Figure 18 Schematic of sampling recommendations.	19

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