

CERTIFICATE OF ANALYSIS

Order Type: Medical Cannabis
Order ID: OR2019-974

Customer ID: 37
Customer Name: Can-Tek Labs

Harvest/Extract Lot: None
Harvest/Extract Batch: None

Cultivar (Strain): Sat a vet 100mg PCR
 Extract Horse

Lab ID: SA2019-3347

Sample Matrix: Oil/Tincture

Sample Date: 04/05/2019

Date Received: 04/05/2019

Date Completed: 04/08/2019

Remarks:

CANNABINOID (POTENCY) PROFILE

Analysis Date/Time: 04/06/2019 0345

Method: HPLC/DAD (Internal Method-001)

Moisture Content (%): -

Analyst: OL

Instrument: Agilent 1100

Water Activity (aw): -

Cannabinoid	Result (%)	Result (mg/g)	Reporting Limit (mg/g)	Per Unit (mg)
CBD	0.0303	0.303	0.0025	36.4
CBDa	-	-	0.0025	-
CBDv	-	-	0.0025	-
Δ9-THC	-	-	0.0025	-
Δ8-THC	-	-	0.0025	-
THCa	-	-	0.0025	-
THCv	-	-	0.0025	-
CBC	-	-	0.0025	-
CBG	-	-	0.0025	-
CBGa	-	-	0.0025	-
CBN	-	-	0.0025	-



TOTAL 0.0303 0.303 36.4

UNIT MASS (g): 120

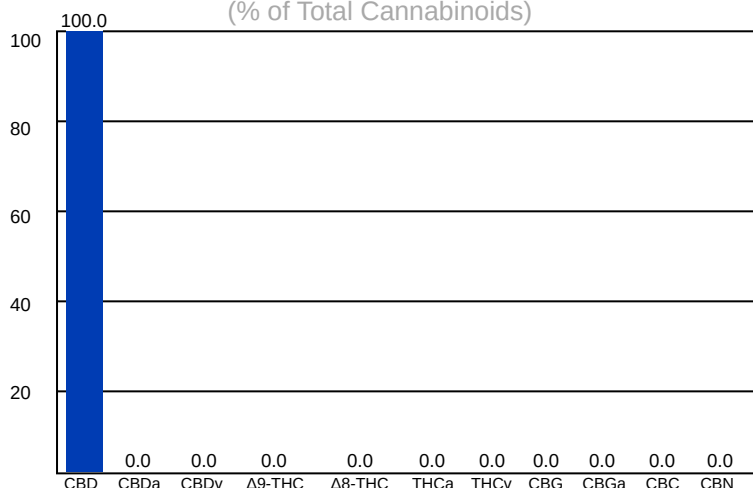
TOTAL THC - - -

"-" Not detected above RL.

TOTAL CBD 0.0303 0.303 36.4

Cannabinoid Distribution

(% of Total Cannabinoids)



Deviations from standard operating procedure: None

Recoveries for all analyte standards: 90-110%

Replicate Uncertainties: <5% RSD, <20% RPD

Sample/Reagent Blanks: <RL for all analytes

Values for plant matter are adjusted for moisture content.

Total THC = (THCa x 0.877) + Δ9-THC

Total CBD = (CBDa x 0.877) + CBD

Percentage results are reported by mass.
 mg/g results are reported as mass component per mass material.

Abbreviations: UV - Ultraviolet, HPLC - High Pressure Liquid Chromatography, RL - Reporting Limit, RPD - Relative Percent Difference, RSD - Relative Standard Deviation

This information is provided as a service and makes no claims of efficacy and/or safety of this product. Results are applicable only for the sample(s) analyzed and for the specific analysis conducted. This report is for informational purposes only and should not be used to diagnose, treat, or prevent any medical-related symptoms.

The statements and results herein have not been approved and/or endorsed by the FDA.

REPORT OF LABORATORY ANALYSIS

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