

Sample Preparation of Cannabis Products by Syringe Filtration using the Shimadzu Cannabis Analyzer

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Summary

Cannabis flower is an extremely complex matrix that must be filtered prior to HPLC or LCMS analysis to avoid clogging of the instrument. We evaluated syringe filters used during the sample preparation portion of the Cannabis Analyzer for Potency. The analysis was conducted using the Shimadzu Cannabis Analyzer for Potency High Sensitivity Method (CAP-HS). This analyzer is the most widely used HPLC instrument for determining the potency of cannabinoids found in Cannabis flower. Seven different syringe filter types were tested to determine which filter type was best for filtration of extracted cannabis flower. Three filter types were found to yield acceptable results but the Polypropylene (PP) filter was best.

Introduction

Many states across the United States are legalizing Cannabis either for medicinal or recreational use. Laboratories testing this material are challenged with providing quality testing with varying reporting limits from state to state in an array of different matrices. This is why Shimadzu has developed three different methods for potency analysis of cannabinoids plus a standardized extraction method.

The extraction method currently being utilized for flower by many laboratories is a liquid-liquid extraction with filtration of the extract by using a 0.45um syringe filter prior to HPLC analysis.

Method

In this study, we evaluated the performance of the Shimadzu SyrFi™ syringe filter products. Since extraction efficiency varies by laboratory and by analyst, we conducted both a solvent spiking and matrix spiking evaluation of the filters used in this method. Using the extraction protocol provided with the Cannabis Analyzer for Potency, four representative cannabis flower samples were obtained, extracted, and homogenized to yield a final sample volume of 120mL. Additionally, Calibrators and Quality Control samples were prepared. Both QC samples were run before and after each filter type.

A 1mL aliquot of the sample was segregated as the unspiked unfiltered sample. The extracted sample was spiked using Shimadzu's 11-part cannabinoid mix (220-91239-21) to a concentration of 10ppm with a final volume of 100mL. 1mL of the spiked sample was segregated as the spiked unfiltered sample. 1 mL volumes of sample were filtered separately through each of the syringe filters in replicates of ten. The 0.45um porosity, 13mm diameter syringe filters were used for filtration.

Each sample, standard and QC was analyzed using the 10-minute CAP-HS method with UV detection. We accomplished near baseline resolution using a Shimadzu NexLeaf C18 2.7µ analytical column (220-91525-70) with associated guard column, as seen in the chromatogram in Figure 1.

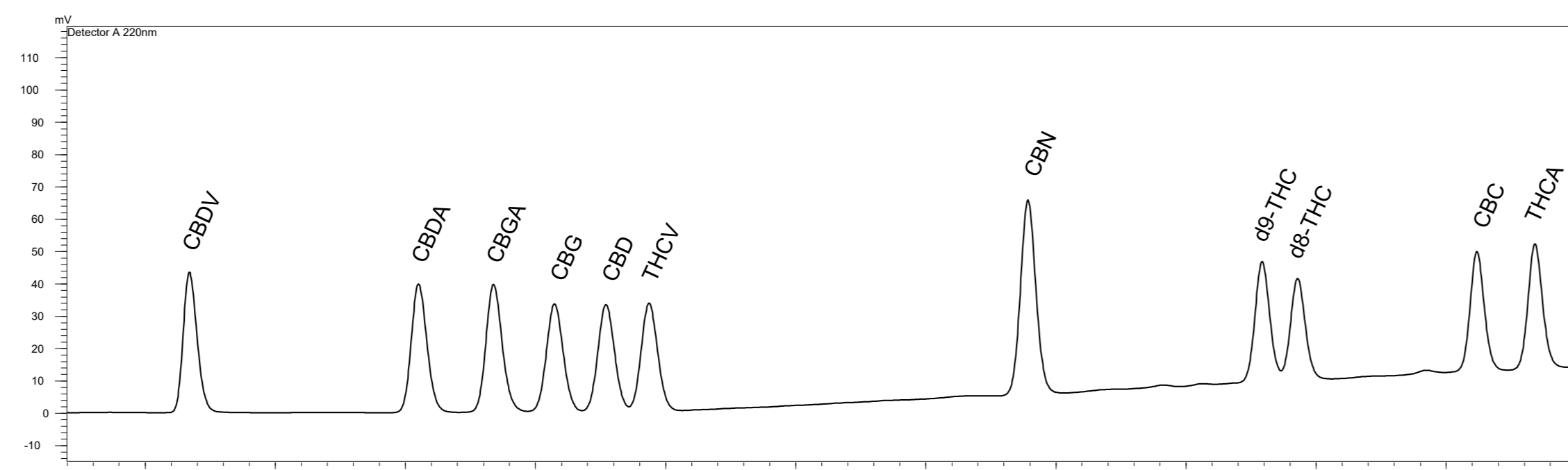


Figure 1: A representative chromatogram of the 10ppm calibration point showing separation of all Cannabinoids.

Results and Discussion

Initial Calibration

A series of six initial calibration standards across the range of 0.5 to 100 ug/mL (parts-per-million, ppm) and two Quality Control (QC) samples, one at 20ppm and one 80ppm were prepared. The calibration curve was evaluated using both correlation coefficient (r2) from a linear regression and using the percent relative standard deviation (% RSD) for each data point in the curve. All calibration curves passed the CAP-HS criteria (RSD < 20%, r2>0.9900). Figure 2 shows the calibration curves for all compounds.

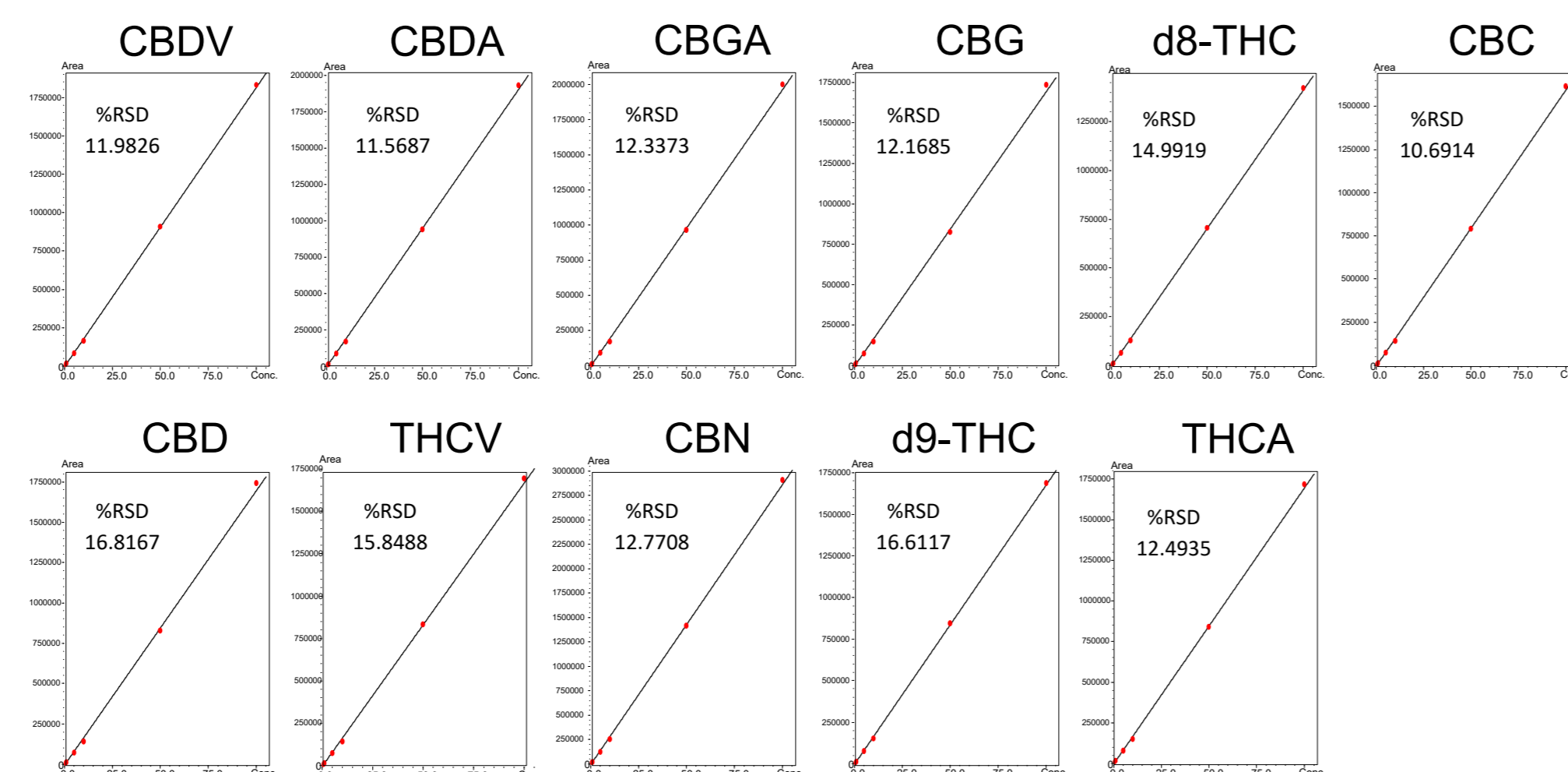


Figure 2: Calibration curves for all compounds contained in CAP-HS

Quality Control Standards

Quality Control (QC) standards with a concentration of 20ppm and 80ppm for all compounds were analyzed before and after each new filter type was tested. The QC concentrations were calculated based on the initial calibration curve, and recoveries were between 70 to 130% which is within the calibration acceptance criteria. Table 1 shows the statistical results for the initial calibration curves and two representative QCs. Figure 3 shows a representative chromatogram of both the high and low QC's.

Compound	Standards			OQ High		OQ Low	
	%Dev	Accuracy[%]	R ²	%RSD	%Dev	Accuracy[%]	%Dev
CBDV	5.010	100.011	0.999	11.983	2.850	97.100	13.32
CBDV	7.361	1.115	6.459	1.843	6.873	1.365	6.691
CBDV	1.115	6.459	1.843	6.873	1.365	6.691	2.408
CBDV	6.873	1.365	6.691	2.408	7.171	4.573	6.873
CBDV	1.441	85.60	6.824	1.371	6.824	1.441	6.824
CBDV	1.371	6.824	1.441	6.824	1.371	6.824	1.441
CBDV	1.441	6.824	1.371	6.824	1.441	6.824	1.371
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