



# Analysis of Cannabinoids in Hemp using QuEChERS Extraction, SpinFiltr™ dSPE Cleanup and LC-MS/MS Detection

## UCT Part Numbers

### ECMSSC-MP

Mylar pouch containing  
4g MgSO<sub>4</sub> and 1g NaCl

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### ECQUSF54CT

SpinFiltr™ dSPE cleanup tube  
150 mg MgSO<sub>4</sub>, 50 mg PSA,  
50 mg C18 and 50 mg ChloroFiltr®

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### SLAQ100ID21-3UM

Selectra® Aqueous C18  
HPLC column  
100 × 2.1 mm, 3 μm

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### SLAQGDC20-3UM

Selectra® Aqueous C18  
guard cartridge  
10 × 2.1 mm, 3 μm

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### SLGRDHLDR

Guard cartridge holder



## Summary:

Thirty-nine states have defined industrial hemp as an agricultural product distinct from cannabis. Nineteen states conducted hemp research in 2017, including Colorado, Hawaii, Indiana, Kentucky, Minnesota, Montana, Nebraska, Nevada, New York, North Dakota, Oregon, Pennsylvania, Tennessee and Washington.

This application note outlines a QuEChERS method for the analysis of hemp for three cannabinoids. Sample purification is carried out using UCT's new cleanup product SpinFiltr™, which combines the convenience of classical dispersive-SPE (dSPE) with an ultrafiltration tube containing a 0.2 μm filter membrane to simultaneously remove unwanted matrix components and filter the sample prior to LC or GC analysis. The SpinFiltr™ dSPE tube uses PSA, C18 and ChloroFiltr® sorbents for sample cleanup. ChloroFiltr® is a unique polymeric sorbent designed for the removal of chlorophyll. Liquid chromatography, using a Selectra® Aqueous C18 column, coupled to tandem mass spectrometry (LC-MS/MS) is used for analysis of the cannabinoids.

Spin  Filtr™



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## Sample Pretreatment:

Hemp should be ground to a fine powder using cryogenic milling (e.g. SPEX 6775 Freezer/Mill®). For this work a large quantity (100 g) of hemp was thoroughly blended in a Robot-Coupe® using dry ice to generate a homogenous sample for use during method development and recovery experiments.



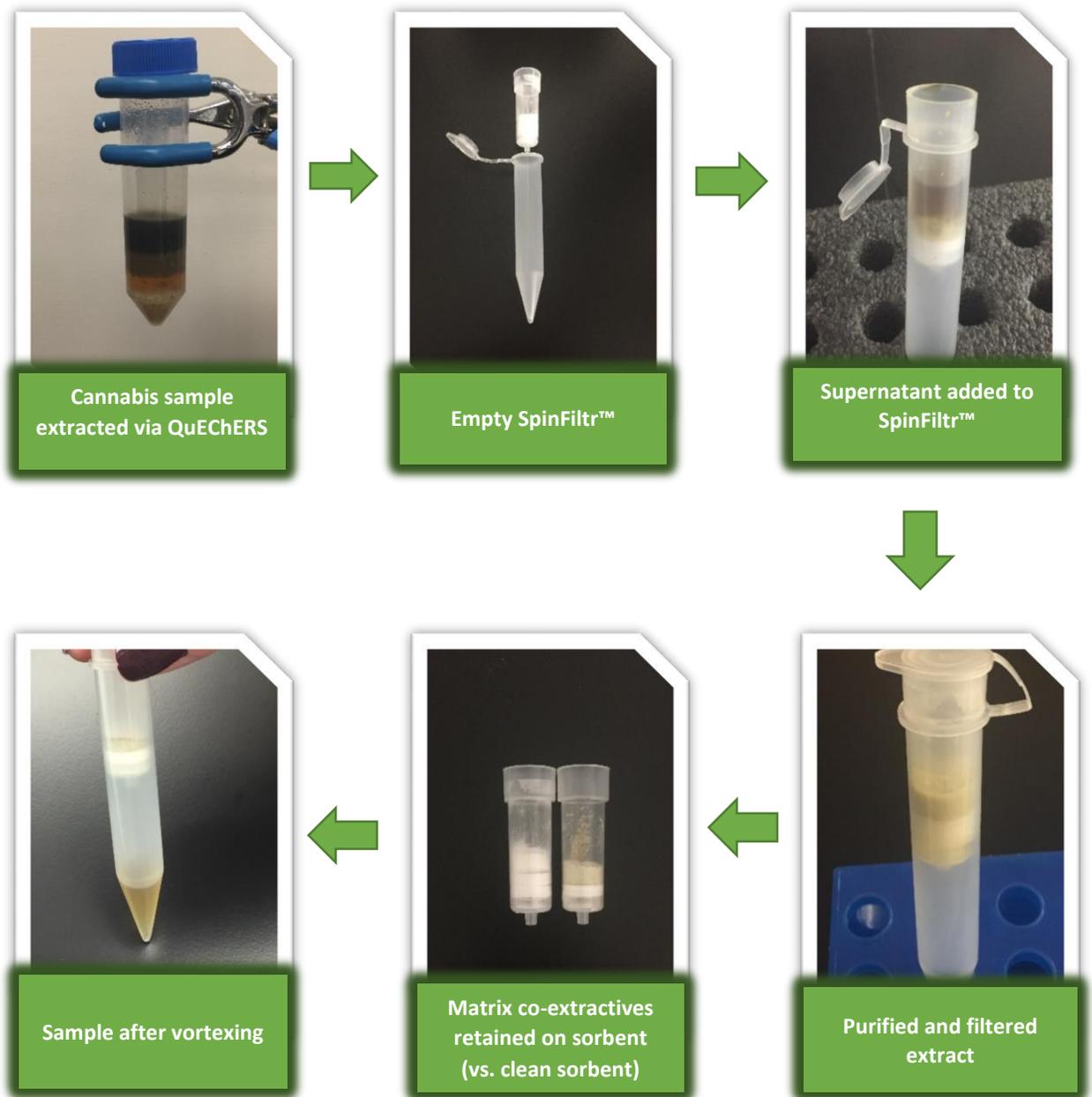
**Figure 1.** Hemp sample before (left) and after (right) homogenization with dry ice.

## QuEChERS Procedure:

1. Weigh 1 g of hemp sample into a 50 mL polypropylene centrifuge tube.
2. Add internal standard(s).
3. Add 10 mL ultrapure water, vortex briefly, and allow sample to hydrate for 15 min (improves extraction efficiency).
4. Add 10 mL acetonitrile containing 2% formic acid.
5. Add the contents of the **ECMSSC-MP** Mylar pouch and shake for a minimum of 5 minutes (by hand or mechanically). For this work a Spex 2010 Geno/Grinder® was used (1500 RPM).
6. Centrifuge the sample at  $\geq 3000 \times g$  for 5 minutes.
7. Transfer 1 ml of supernatant to a SpinFiltr™ dSPE cleanup tube (**ECQUSF54CT**).
8. Vortex the sample for 30 seconds.
9. Centrifuge the sample at  $\geq 3000 \times g$  for 5 minutes.
10. Transfer the purified and filtered sample extract into an autosampler vial for analysis. \*

\*Note: LC-UV can also be used for final cannabinoid content analysis





**Figure 2.** Schematic of the SpinFiltr™ dSPE cleanup procedure.



## LC-MS/MS Parameters:

**Table 1. Instrumentation**

<b>UHPLC system</b>	Thermo Scientific™ Dionex™ Ultimate™ 3000
<b>MS system</b>	Thermo Scientific™ TSQ Vantage™ (MS/MS)
<b>HPLC column</b>	UCT Selectra® Aqueous C18, 100 × 2.1 mm, 3 μm (p/n: SLAQ100ID21-3UM)
<b>Guard column</b>	UCT Selectra® Aqueous C18, 10 × 2.1 mm, 3 μm (p/n: SLAQGDC20-3UM)
<b>Guard column holder</b>	p/n: SLGRDHDR
<b>Column temperature</b>	40°C
<b>Flow rate</b>	300 μL/min
<b>Injection volume</b>	5 μL

**Table 2. Mobile Phase Gradient**

<b>Time (min)</b>	<b>% Mobile Phase A (Water + 10mM NH<sub>4</sub>HCO<sub>2</sub>)</b>	<b>% Mobile Phase B (Methanol + 0.1% formic acid)</b>
0.0	40	60
0.5	40	60
3.0	5	95
7.0	5	95
7.1	40	60
10.0	40	60

**Table 3. MS Parameters and Retention Times**

<b>Analyte</b>	<b>RT</b>	<b>Parent ion</b>	<b>Product 1</b>	<b>CE 1</b>	<b>Product 2</b>	<b>CE 2</b>
CBD	5.26	315.0	193.1	20	123.0	30
CBN	5.75	311.1	223.1	19	293.2	14
THC	5.91	315.2	193.1	19	123.1	31

### Conclusion:

The method outlined in this application note allows for the simultaneous analysis of three cannabinoids in hemp in one simple QuEChERS extraction procedure, thereby saving time, sample and cost. Sample cleanup is carried out by dSPE using UCT's new Spinfilter™ product which purifies and filters the sample in one easy step. Chlorofiltr® dSPE sorbent was used to selectively remove chlorophyll. Analysis of the samples was performed by LC-MS/MS utilizing a Selectra® Aqueous C18 HPLC column. With the increase of hemp usage across the United States, this simple method will be beneficial for any research facility wanting to ensure the cannabinoid content in their product.

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