AN092



Cannabis Potency analysis by GC-FID with hydrogen carrier gas

Scion – Advanced Projects & Custom Solutions



Introduction

Hemp and marijuana are becoming more popular because of legalization in multiple countries. The increase of this market also increases the offer of products that contain very different levels of cannabinoids than the content described on the label. Therefore, as quality control it is important to monitor the level of cannabinoids.

Marijuana usually contains high levels of $\Delta 9$ -THC and a lower level of CBD, with hemp it is the other way around. The primary psychoactive component is $\Delta 9$ -THC, while CBD is the primary therapeutic component.

Consumer hemp generally comes in the form of hemp oil which is used for medical purposes while marijuana is often smoked. Keep in mind that both samples need different sample preparation before analysis due to the difference in sample matrixes.

Potency analysis identifies different cannabinoids and measures their concentration as an indication of the strength of the product. There are over 500 chemical compounds in Cannabis, the most tested cannabinoids can be found in table 1. These are also discussed in this application note.

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Component	Comment
CBC (Cannabichromene)	
CBD (Cannabidiol)	
Δ8-THC (Δ8-tetrahydrocannabinol)	
Δ9-THC (Δ9-tetrahydrocannabinol)	
CBG (Cannabigerol)	
CBN (Cannabinol)	
CBDV (Cannabidivarin)	Not detectable by GC unless the sample is derivatised.
CBGA (Cannabigerolic acid)	Not detectable by GC unless the sample is derivatised.
THC-A (Δ9- tetrahydrocannabiolic acid)	Not detectable by GC unless the sample is derivatised.
CBDA (Cannabiolic acid)	Not detectable by GC unless

the sample is derivatised.

Injector	Splitless 50:1, 275 °C
Column	Scion-35MS
Oven Program	150°C (0.1 min), 25°C/min to 310°C (0.1 min)
Carrier	Hydrogen, 1.6 ml/min (integrated hydrogen safety bundle)
Detector	FID with ceramic jet, 300°C Air: 300 ml/min, Fuel gas (H ₂): 30 ml/min, Make up (N ₂): 30 ml/min
Inj. Volume	0.5 μΙ
Autosampler	8400
Software	Compass CDS

Table 2. Analytical conditions

Experimental

This analysis can be implemented on both the 436-GC as well as on the 456-GC platform. The Scion 436-GC analyser used for this application is equipped with an 8400 autosampler.

The analysis of these components can be done using a GC with a FID detector, but it is also possible to use a Mass Spectrometer (MS). A different method is using a HPLC, in this case it is also possible to detect THC-A, CBDA, CBDV and CBGA with extra sample preparation (derivatization).

The focus of the calibration lies on six components: CBC, CBD, Δ8-THC, Δ9-THC, CBG and CBN. Unfortunately, CBDV, CBGA, THC-A and CBDA cannot be detected without extra sample preparation due to decarboxylation caused by the heat of the injector. Decarboxylation is the removal of a carboxyl group releasing carbon dioxide. CBDV decarboxylates to CBD, CBGA to CBG, THC-A to THC and CBDA to CBD.

The Quality Control (QC) sample was made from 20 μ g/ml for all components.

Different products were examined to determine potency. Cannabis oil and medical marijuana were chosen in this study.



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The cannabis oil was diluted with methanol, for the medical marijuana a bit more preparation was necessary. In order to extract the potency from the medical marijuana, it had to be dried for 2 hours, after that it had to be grinded. 30 ml of methanol was added and placed in an ultrasonic bath for 30 minutes. Evaporate to dryness, reconstitute in methanol.

When looking at the settings of the GC it is important that the injector temperature is high enough to prevent carry over. Carry over can occur from previous runs or extra material that is being introduced into the system.

The chromatogram in figure 1 shows carry over from the previous run. When the injector temperature was increased it showed that the carry over disappeared (see figure 2). Another solution for this problem could be a washing method between samples.

Both figure 1 and 2 are examples from Scion Instruments application note AN091 which explains the analyses of potency using a helium carrier gas instead of hydrogen.

In this application note 150 °C is used as the starting temperature developing this method. The total time with this starting temperature is 6.6 minutes, it is possible to speed-up the method.

Figure 3 shows the four chromatograms with the possible starting temperatures, 225 °C will be the fastest method. Figure 3 however, shows a contamination peak between 9-THC and CBG. When the starting temperature is increased the separation of this contamination peak and 9-THC will decrease.

150 °C as starting temperature provides the best results with minimum interference. When choosing a faster method, it is advised to keep a good eye on this disturbances that is not the wanted analyte.

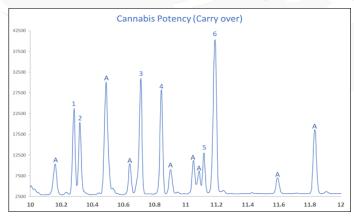


Figure 1. Chromatogram of a potency standard showing carry over (A) using helium carrier gas (AN0091)

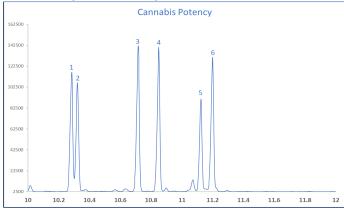


Figure 2. Chromatogram of a potency standard using helium carrier gas (AN0091)

Peak ID	Compound Name	
1	CBC*	
2	CBD*	
3	8-THC	
4	9-THC	
5	CBG	
6	CBN	
Α	Carry over peak	

*With helium carrier gas 1 is CBD and 2 is CBC, the switch of the components is caused by using a 5-ms column instead of a 35MS

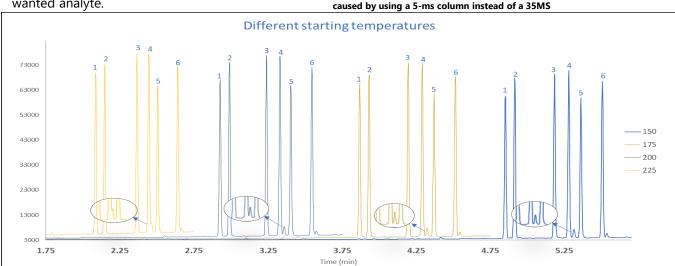


Figure 3. Chromatograms with different starting temperatures using a hydrogen carrier gas

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Results

The precision of the method was obtained by ten consecutive injections containing 50 μ g/ml of each component. Most standards described that the RSD% of an accurate method must be no more than 2%, CBC showed an RSD of 1.2%, CBD 1.0%, Δ 8-THC 1.0%, Δ 9-THC 1.1%, CBG 1.9% and CBN 1.11%. The precision performance of this method is excellent

When looking at the chromatogram of figure 3 it is shown that all components are base line separated, this means accurate quantitation can take place.

In addition to these results, it is shown that the peak tailing factor (PTF) of all components is smaller than 2. CBD 0.95, CBC 0.96, 8-THC 1.03, 9-THC 1.02, CBG 1.06 and CBN 1.01. This overall is an outstanding result.

The calibration curves for the potency standards were prepared between the 2 to $100 \mu g/ml$. Figure 4 shows the calibration range for all the standards and it showed that all the correlation coefficients (R²) bigger than 0.999 which is exceptionally good.

Originally the linearity the limit of detection (LOD) and limit of quantitation (LOQ) was calculated, these results can be found in table 3.

Component	LOD (µg/ml)	LOQ (µg/ml)
CBC	1.0	3.1
CBD	0.8	2.3
8-THC	0.72	2.2
9-THC	0.73	2.2
CBG	0.91	2.8
CBN	0.97	2.9

Table 3. LOD and LOQ for the different components

The QC sample was analysed using ten injections. The average concentration for CBD was 18.9 μ g/ml with an RSD% of 0.89, CBC was 18.4 μ g/ml with an RSD% of 0.54. 8-THC was 20.8 μ g/ml with an RSD% of 0.9 , 9-THC was 20.5 μ g/ml with an RSD% of 0.86, CBG was 16.9 with an RSD% 2.87 and CBN was 20.3 with an RSD% of 0.99.

The analysis of two cannabis samples was performed on an oil and a medical marijuana sample.

According to the label of the oil sample it should contain 100 mg of CBD, other components were not mentioned on the label.

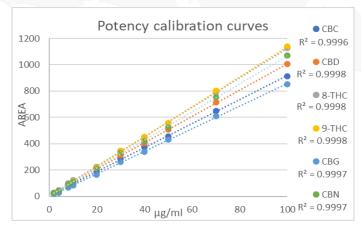


Figure 4. Calibration curves of the potency standards

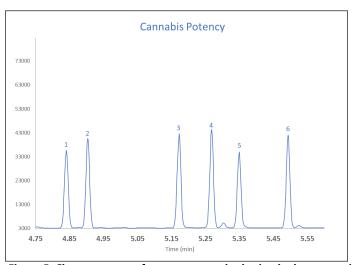


Figure 5. Chromatogram of a potency standard using hydrogen carrier.

Peak ID	Compound Name	
1	CBC	
2	CBD	
3	8-THC	
4	9-THC	
5	CBG	
6	CBN	

After analysis it showed that the oil sample contained 65.3 mg CBD, CBC and 9-THC were also detected, unfortunately these were below the LOQ, it also shows that 8-THC, CBG and CBN were not present in this sample.

The medical marijuana did not come with an index stating it's content. Analyses showed it contained 12.7% CBD and 0.79% 9-THC. CBC, 8-THC and CBG were detected but the concentration was below the LOQ,. CBN was not present in this sample.

Figure 6 shows an example chromatogram of the oil and medical marijuana sample.



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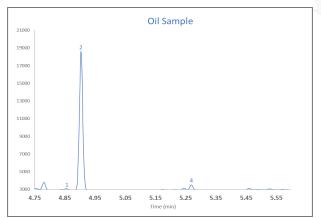
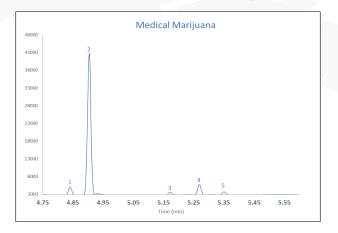


Figure 6. Chromatogram of an oil and medical marijuana sample

Conclusion

The Scion 4X6-GC analyser equipped with a split/spitless injector, scion instruments column and FID is capable of analysing potency from cannabis products in a qualitative and quantitative way.

Although the method developed is well suited for potency analysis it is confirmed that one of the disadvantages of this method is that the acidic form of the potency components is converted in the hot injector. It should be kept in mind that the levels of CBD, CBG and 9-THC in this case is a summation of the non-acidic and acidic form. If the acidic form has to be analysed it is advised to perform extra sample preparation or another analysing method in the form of a Scion Instruments 6000 HPLC method can be chosen.



Peak ID	Compound Name
1	CBC
2	CBD
3	8-THC
4	9-THC
5	CBG
6	CBN



Picture 1: The Scion 4X6-GC series with 8400 autosampler

SCION Instruments

UK

Livingston Business Centre Kirkton South Road, Livingston West Lothian EH54 7FA Scotland, UK Phone +44 1506 300 200 sales-eu@scioninstruments.com

The Netherlands

Amundsenweg 22-24 4462 GP Goes, The Netherlands Phone +31 (0) 113 287 600 sales-eu@scioninstruments.com