

# Alcohols $C_1$ - $C_3$ on an Agilent J&W PoraPLOT Q GC Column

## **Application Note**

Forensic Toxicology

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#### Introduction

The conversion of methanol into ethanol by reaction with hydrogen and carbon monoxide is well known. This reaction is carried out in the presence of a water-soluble cobalt catalyst at elevated temperatures and pressures. Higher alcohols are formed in relatively small amounts as by-products.

 $C_1$ - $C_3$  alcohols are polar solvents and are analyzed in a wide variety of samples including blood for alcohol intoxication. This application note shows the separation of these alcohols on an Agilent J&W PoraPLOT  $\Omega$  column.



## **Materials and Methods**

Technique: GC-FID

Column: Agilent J&W PoraPLOT Q,

 $25 \text{ m} \times 0.32 \text{ mm df} = 10 \mu\text{m} (p/n CP7551)$ 

Sample: Compounds in headspace

Injection Volume: 1 μL

Carrier Gas: Hydrogen, constant pressure, 100 kPa

(1 bar, 14.5 psi)

Temperature: 200 °C

Injection: 275 °C, split 1:100

Detection: FID, 275 °C

## **Results and Discussion**

The analysis of the  $\rm C_1$ - $\rm C_3$  alcohols took less than 2 minutes with the PoraPLOT Q column. The compounds were baseline separated and had an excellent peak shape (Figure 1).

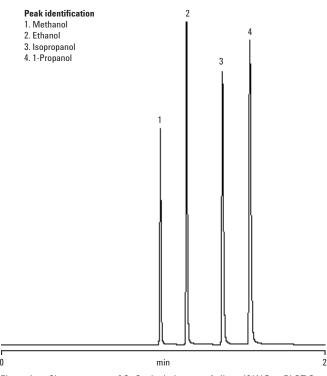


Figure 1. Chromatogram of  $C_i$ - $C_s$  alcohols on an Agilent J&W PoraPLOT Q column.

## **Conclusion**

Using a PoraPLOT  $\Omega$  GC column, separation of  $C_1$ - $C_3$  alcohols was achieved in less than 2 minutes. PoraPLOT  $\Omega$  is recommended for column switching systems that analyze polar and apolar volatile compounds. The column delivers repeatable retention times because retention is not influenced by water in the sample.

### Reference

http://www.freepatentsonline.com/4424383.pdf

#### www.agilent.com/chem

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