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Successful Separation of Delta-8-Carboxy-THC and Delta-9-Carboxy-THC by LC-MS/MS Analysis

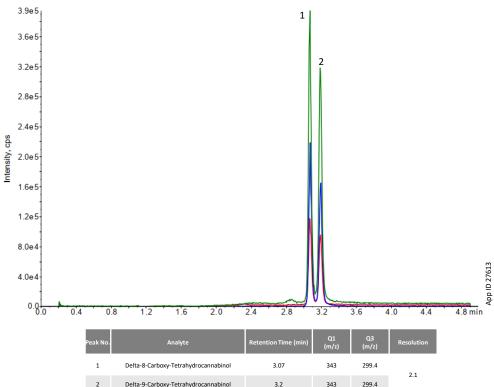
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Overview

Delta-8-Tetrahydrocannabinol (delta-8-THC) is a structural isomer to Delta-9-THC and produces similar mind-altering affects. It is sold as an alternative to marijuana which, while legal in many states, is a DEA Schedule I drug and still illegal under federal law. Legal status of Delta-8-THC is murky and varies by jurisdiction. This has led to an increase in availability and use of Delta-8-THC by drug users looking for a "legal" high. Distinguishing use of Delta-8-THC from Delta-9-THC is important for forensic laboratories conducting drug testing.

Urine is a common specimen used for forensic drugs of abuse testing and workplace drug testing. "Parent" Delta-8-THC and Delta-9-THC are extensively metabolized to their respective carboxy metabolites. The two analytes have similar structures and the same mass (and hence the same m/z), so these two metabolites need to be chromatographically separated for accurate LC-MS/MS identification and quantitation.

This application note provides conditions for separation of the carboxy metabolites of Delta-8-THC and Delta-9-THC using a Synergi 2.5 μ m Hydro RP column.





LC Conditions

Column: Synergi™ 2.5 μm Hydro RP

Dimensions: 100 x 3.0 mm **Part No.:** 00D-4387-Y0

Mobile Phase: A: 5 mM Ammonium Formate

B: Methanol

Gradient: Time (min) %B
0 70
1 95
3.5 95
3.51 70
5 70

Flow Rate: 0.4 mL/minInjection Volume: $10 \text{ }\mu\text{L}$ Temperature: $25 \text{ }^{\circ}\text{C}$

LC System: Agilent® 1260 Infinity

Detection: MS/MS

Detector: SCIEX® 6500 Triple Quad™

Analyte Concentrations: 10 ng/mL

MS/MS Conditions

Ion Source:	ESI
Polarity:	Negative
Temperature:	500 °C
GS1:	60 psi
GS2:	50 psi
CUR:	35 psi
IS:	-3500 V
CAD:	Medium
DP:	-40 V
EP:	-10 V
CE:	-35 V
CXP:	-13 V

Need a different column size or sample preparation format?

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