

# MS/MS Identification of Four Aflatoxins Using the Agilent 500 Ion Trap LC/MS

## **Application Note**

Food Testing and Agriculture

## Abstract

A rapid and sensitive LC/MS/MS method for the detection and analysis of aflatoxins B1, B2, G1 and G2 provides detection down to 1 ppb. The 500 Ion Trap MS/MS capabilities result in the collection of clear, baseline resolved chromatographic peaks for target compound quantitation.

## Introduction

Aflatoxins are a group of structurally related carcinogenic mycotoxins produced by *Aspergillus sps*. The most commonly found aflatoxins in food and feed are aflatoxins B1, B2, G1 and G2. Aflatoxin B1 has been classified as a Class 1 human carcinogen by the International Agency for Research on Cancer. Globally, there are significant regulations on the presence of aflatoxins in food and feed. The U.S. Federal Government action level for aflatoxins in food for human consumption or in dairy cow feed is 20 ppb.

The European Commission has set very low action levels for aflatoxins in cereals intended for direct human consumption or for ingredients in foodstuffs. These action levels are 4 ppb total aflatoxins and 2 ppb for aflatoxin B1. So far, these compounds have been extensively analyzed by HPLC. This study developed a rapid and sensitive LC/MS/MS method for the detection and analysis of aflatoxins B1, B2, G1 and G2.



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#### Instrumentation

The following instruments were used in this study:

- Agilent 500 Ion Trap LC/MS with ESI source
- Agilent 212-LC Binary Gradient LC/MS pumps
- Agilent Prostar 430 AutoSampler

#### **Materials and Reagents**

The aflatoxins used were purchased from Supelco, Bellefonte, PA. All other reagents were purchased from Sigma-Aldrich Co, St. Louis, MO.

#### **HPLC** conditions

Column:	Polaris C18-A, 150 × 3 mm, 5 µm (Agilent p/n A2000150X030)					
Solvent A:	Water					
Solvent B:	Methanol					
Injection volume:	10 µL					
LC program:	Time			Flow		
	(min:sec)	%A	%В	(µL/min)		
	00:00	70	30	200		
	01:00	70	30	200		
	15:00	10	90	200		
	15:01	0	100	200		
	20:00	0	100	200		
	20:01	70	30	200		
	25:00	70	30	200		

#### **MS** parameters

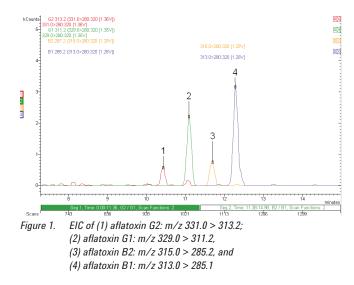
Ionization mode:	ESI Positive		
API drying gas:	35 psi at 350 °C		
API nebulizing gas:	40 psi		
Needle:	5000 V		
Shield:	600 V		

#### Table 1. MS Segment Parameters

Segment	Analyte	Transition	Retention time (min)	Capillary voltage (V)	Excitation amplitude (V)	RF load %
1	Aflatoxin G2	331 → 280–320	10.45	75	1.36	82
1	Aflatoxin G1	329 → 280–320	11.12	75	1.35	82
2	Aflatoxin B2	315 → 280–320	11.68	75	1.29	82
2	Aflatoxin B1	313.2 → 280–320	12.30	75	1.28	82

#### **Results and Discussion**

This LC/MS/MS method separates aflatoxins G2, G1, B2 and B1, and provides excellent response for each of the analytes down to 1 ppb. The Polaris C18 column provides excellent chromatographic separation and reproducible retention times. Figure 1 shows the extracted ion chromatogram (EIC) of aflatoxin G2, G1, B2 and B1 at a concentration of 1 ppb.



Calibration curves were found to be linear from 1 to 50 ppb, with %RSD values of 15.62% for aflatoxin G2, 13.64% for aflatoxin G1, 12.65% for aflatoxin B2 and 10.44% for aflatoxin B1. Figure 2 shows the calibration curve for aflatoxin G1.

Calibration Curve Report File: ...atoxins\aflatoxin\_02-msms 2seg@500,000 200ul 1uscan@1hz cal\_2.mth Detector: 500-MS Mass Spec, Address: 56

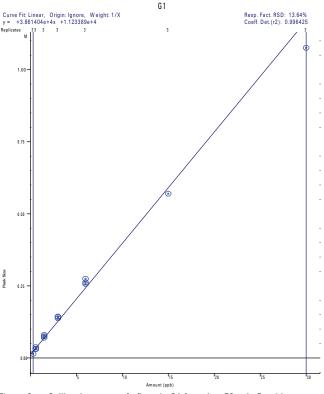


Figure 2. Calibration curve of aflatoxin G1 from 1 to 50 ppb. For this curve, %RSD is 13.64% and  $r^2 = 0.996$ .

## Conclusion

The four aflatoxins presented in this method were separated and identified in less than 15 minutes.

The MS/MS capabilities of the Agilent 500 Ion Trap LC/MS allow for isolation of desired precursor ions followed by collision induced dissociation (CID) and characteristic product ion spectra, resulting in the collection of clear, baseline resolved chromatographic peaks for target compound quantitation. This method is fast, rugged and sensitive.

#### References

- 1. Commission regulation (EC) No 2174/2003 / (EC) No 466/2001 as regards aflatoxins.
- 2. U.S. Food and Drug Administration Industry Activities Staff Booklet August 2000.
- 3. European Commission Council directive 96/23/EC/SANCO/1805/2000.

## **For More Information**

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