Mycotoxins: Sample Preparation and Analysis

Matrix of the Month

April, 2013:

Aflatoxins in Olive Oil



Do you have a special matrix that we should test for mycotoxins? Please let us know and write an e-mail to info@LCTech.de!

Protocol

10 g sample are mixed with 2 g sodium chloride.

100 mL 80/20 methanol/water and 50 mL n-hexane are added, stir for 30 minutes.

The sample is freezed to -20°C, fat and n-hexane are completely removed, the sample is once more centrifugated at 1500 for 10 minutes.

The lower phase is used (2 mL + 12 mL PBS/Tween (8 %)). 14 mL are added onto the immunoaffinity column AflaCLEAN and then the sample flask and the column are washed with 10 mL deionised water.

The column is dryed and eluated with 2 x 1 mL methanol.

The first milliliter methanol should incubate to the column bed for 5 minutes.

The eluate is diluted to the conditons of the HPLC mobile phase and a maximum of 100 mL are injected.

HPLC Conditions

HPLC: Dionex Ultimate 3000, isocratic

Column oven: 36 °C

Separation column: Mycotoxin HPLC column with guard

Flow rate: 1.2 mL/min (water/methanol/acetonitrile(60/30/15(v/v/v))

Fluorescence detection with post column derivatisation (photochemical with UVE)

Excitation wavelength: 365 nm Emission wavelength: 460 nm

Recovery Rates

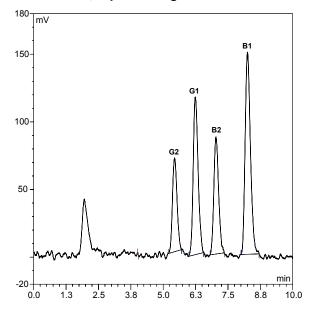
Contents of Aflatoxins B1, B2, G1 and G2 in Olive Oil				
Aflatoxin	B1	B2	G1	G2
Standard*	100	100	100	100
Recovery rate** Olive oil 10 ppb	104	107	90	68

^{*} Standard is set = 100 % , ** corrected with non-spiked sample

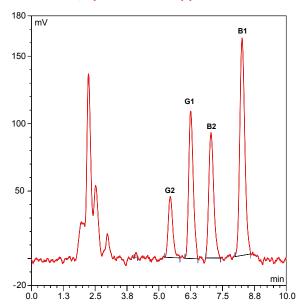


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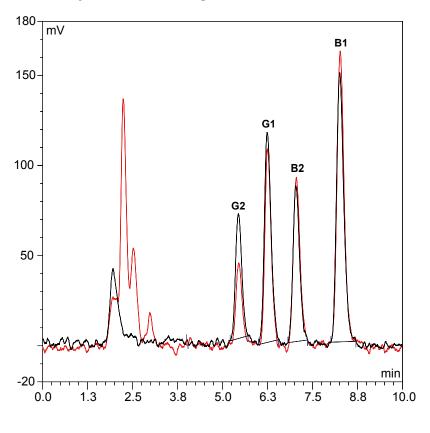
Standard, representing 100 %



Olive oil, spiked with 10 ppb total toxin



Overlay of both chromatograms



These LCTech products were used:

AflaCLEAN,

Immunoaffinity column for the Aflatoxins B1, B2, G1, G2

P/N 10514

UVE,

Photochemical reactor for the analysis of Aflatoxins

P/N 10519

HPLC column,

for the analysis of Aflatoxins

P/N 10522

Do you have further questions? Please simple write an e-mail to info@LCTech.de!