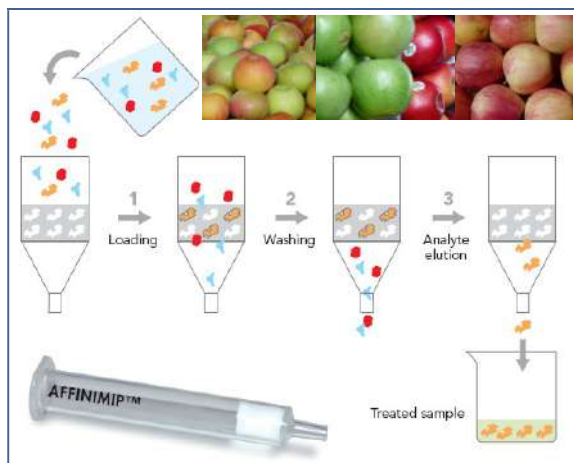


Selective Solid Phase Extraction of Patulin from Apple products Using Molecularly Imprinted Polymers



Introduction

Patulin [4-hydroxy-4*H*-furo[3,2-*c*]pyran-2(6*H*)-one] is a mycotoxin produced by a variety of molds, particularly *Aspergillus* and *Penicillium* species (see figure 1). It is commonly found in rotting apples, and the amount of patulin in apple products is generally viewed as a measure of the quality of the apples used in production.

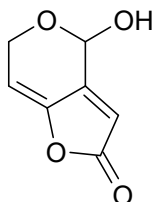


Figure 1. Chemical structure of Patulin, CAS N° 149-29-1

Studies have shown that it is genotoxic. Several countries have instituted patulin restrictions in apple products. Member countries of the European Union have set maximum allowable levels of patulin at 50µg/kg in fruit juices, 25µg/kg in solid apple products, including apple compote, apple puree intended for direct consumption and 10µg/kg in apple juice and solid apple products, including apple compote and apple puree, for infants and young children and in baby foods (European Commission Regulation (EC) 1881/2006 [1]).

Several analytical methods for the determination of Patulin have been developed in which a clean-up step is necessary and crucial. However, by the use of the classical methods of clean-up, the main matrix interferent, 5-Hydroxymethylfurfural (HMF), is still present at a very high concentration, preventing a reliable quantitative Patulin determination.

So there is an increasing need to improve both sensitivity and specificity of this key step of clean-up.

To propose an accurate solution, we have developed a new class of intelligent polymers based on molecularly imprinted polymers specific to Patulin. Molecularly Imprinted Polymer (MIP) is a synthetic material with artificially generated three-dimensional network able to specifically rebind a target molecule. MIP has the advantages to be not only highly selective and specific but also chemically and thermally stable, compatible with all solvents and cost-effective. This polymer is used as a powerful technique for clean-up and pre-concentration applications of Patulin. This study describes the solid phase extraction of Patulin from apple juice and apple puree using a Molecularly Imprinted Polymer (MIP) SPE cartridge that is specific for Patulin (**AFFINIMIP® SPE Patulin**).

Experimental conditions for Apple Juice

Materials

All reagents and chemicals were ACS grade quality or better. Patulin was obtained from Sigma Aldrich (Fluka). Apple juice and apple puree were purchased in different supermarkets.

Solid phase extraction (SPE) protocol for apple juice

The SPE procedure used a 3mL **AFFINIMIP® SPE Patulin** Cartridge. The details of each step are as follows:

- Condition the SPE Cartridge with 2mL of Acetonitrile (ACN), then with 1mL of deionized water
- Load 4mL of the loading solution
- Wash the cartridge with 1mL of NaHCO₃ 1% in water
- Wash cartridge with 2mL of deionized water
- **Force the water down into the cartridge and out the bottom or apply vacuum 10 seconds**
- Wash the cartridge with 1mL of diethyl ether
- Elute Patulin with 2mL of ethyl acetate

The SPE procedure lasted approximately 20 minutes. The elution fraction was then evaporated and dissolved in water containing 0.1% acetic acid. The evaporation time of the elution fraction is approximately 10 minutes.

Analysis

HPLC was performed on a ThermoFinnigan Spectra System with an Atlantis T3 column 150mm x 2.1mm (Waters). The separation was carried out using a mobile phase of deionized water/ACN (95/5, v/v) at a flow rate of 0.2mL/min. The detection system was a ThermoFinnigan Spectra System Model UV6000LP set to 276nm. The injection volume was 100µL.

Results

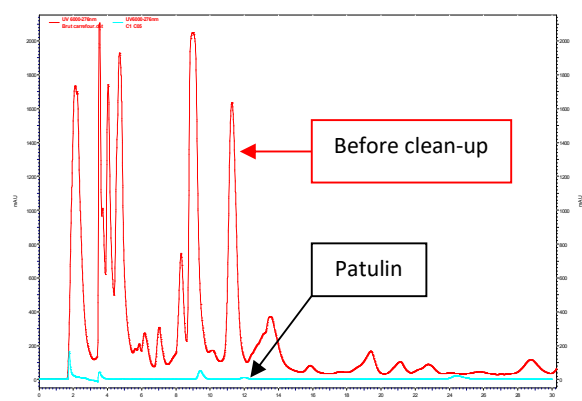


Figure 2. Chromatograms of apple juice containing 25µg/kg of Patulin before (Red) and after (Blue) **AFFINIMIP® SPE Patulin** Clean-up

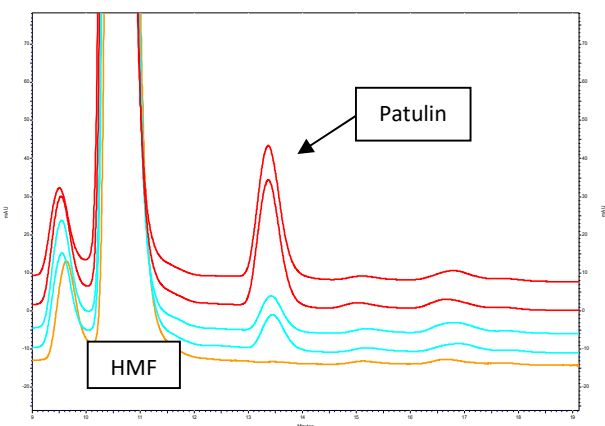


Figure 3. Chromatograms obtained after **AFFINIMIP® SPE Patulin** Clean-up of an apple juice spiked at 40µg/kg (tested twice, red) or at 10µg/kg (tested twice, blue) with Patulin or not spiked (orange)

Table 1. Recovery of Patulin at a contamination level of 10µg/kg and 40µg/kg in apple juice after **AFFINIMIP® SPE Patulin** Clean-up and relative standard deviation calculated from results generated under reproducibility conditions.

Concentration of Patulin (ng/mL)	Recoveries %	% RSD _R
10	97.9	11 (n=9)
40	90.6	11 (n=41)

Experimental conditions for Apple puree

Preparation of samples prior to SPE with **AFFINIMIP® SPE Patulin** Cartridge

Weigh 10g of apple puree; add 150µL of a pectinase enzyme solution followed by 10mL water and mix. Leave solution at room temperature overnight or for 2h at 40°C. Centrifuge at 4500g for 5min and then filter the solution with a 0.2µm filter. This solution is used as the loading solution.

Solid phase extraction (SPE) protocol for apple puree

The SPE procedure used a 3mL **AFFINIMIP® SPE Patulin** Cartridge. The details of each step are as follows:

- Condition the SPE Cartridge with 2mL of acetonitrile (ACN), then with 1mL of deionized water
- Load 5mL of the loading solution
- Wash the cartridge with 4mL of deionized water containing 1% of acetic acid
- Wash the cartridge with 4mL of deionized water
- **Force the water down into the cartridge and out the bottom or apply vacuum 10 seconds**
- Wash the cartridge with 1mL of CHCl₃ (or 500µL of diethyl ether)
- Elute Patulin with 2mL of ACN containing 1% acetic acid (Elution solution may be replaced by 2mL of Ethyl acetate)

The SPE procedure lasted approximately 30 minutes. The elution fraction was then evaporated and dissolved in water containing 0.1% acetic acid.

Analysis

HPLC was performed on a ThermoFinnigan Spectra System with an Atlantis T3 column 150mm x 2.1mm (Waters). The separation was carried out using a gradient (see table 2) at a flow rate of 0.2mL/min:

Table 2. Gradient for the analysis of Patulin after an extraction from an apple puree.

Time (min)	% water	% ACN
0	98	2
20	98	2
21	50	50
25	50	50
26	98	2

The detection system was a ThermoFinnigan Spectra System Model UV6000LP set to 276nm. The injection volume was 100µL.

Results

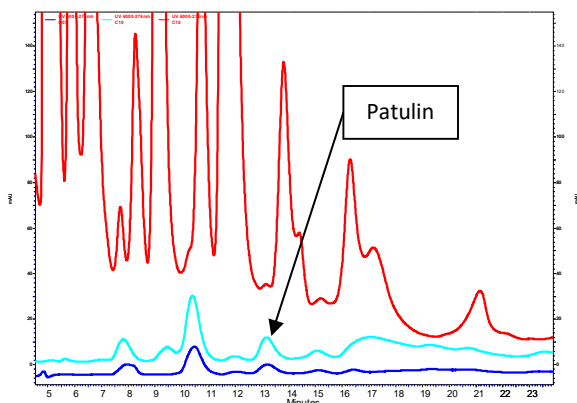


Figure 4. Chromatograms of apple puree containing 40µg/kg or 80µg/kg of Patulin before (Red) and after (Blue) **AFFINIMIP® SPE Patulin** Clean-up

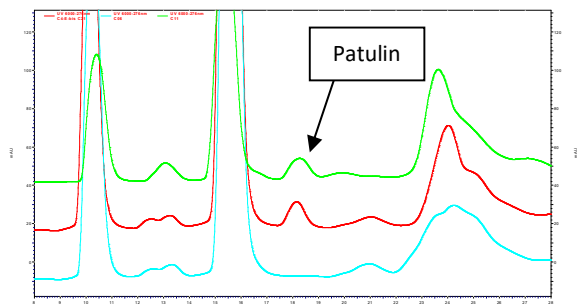


Figure 5. Chromatograms of apple puree containing 0µg/kg (blue) or 20µg/kg (tested twice, green and red) of Patulin after **AFFINIMIP® SPE Patulin** Clean-up

Table 3. Recovery and repeatability of Patulin (n=3) at a contamination level of 20µg/kg in apple puree after **AFFINIMIP® SPE Patulin** Clean-up.

Concentration of Patulin (µg/kg)	Recoveries % (n=3)	% RSDr
20	84	4.5

Conclusion

The use of an **AFFINIMIP® SPE Patulin** cartridge is a simple, fast, sensitive and selective tool for the extraction of Patulin from apple products.

This method complies with the performance criteria for Patulin established by the European Commission Regulation (EC) 401/2006 [2]. This regulation requires recovery values for Patulin higher than 70% for analysis done between 20 to 50µg/kg and higher than 50% for analysis done below 20µg/kg.

The use of **AFFINIMIP® SPE Patulin** enables to obtain recoveries above 80%. This method is well-suited for the analysis of Patulin in apple products.

References

- [1] Commission Regulation (EC) No. 1881/2006 of 19 December 2006, Official Journal of the European Union.
- [2] Commission Regulation (EC) No. 401/2006 of 23 February 2006, Official Journal of the European Union.

Related products

- **AFFINIMIP® SPE Patulin**

Catalog number: FS102-02 for 25 columns

- **AFFINIMIP® SPE Ochratoxin A**

Catalog number: FS101-02 for 25 columns

Please follow the link to [Application note AFFINIMIP® Ochratoxin A Automated protocol \(GILSON GX271 ASPEC™\)](#)

- **AFFINIMIP® SPE Zearalenone**

Catalog number: FS100-02 for 25 columns

Please follow the link to [Application note AFFINIMIP® Zearalenone – Automated protocol \(GILSON GX271 ASPEC™\)](#)