Co-occurrence of aflatoxins and ochratoxin A in spices commercialized in Italy

Ambra Prelle¹, Davide Spadaro¹,², Angelo Garibaldi¹, Maria Lodovica Gullino¹,²

¹ Centre of Competence for the Innovation in the Agro-environmental Sector (AGROINNOVA), University of Turin, Grugliasco (TO), Italy.

² DISAFA, Dept. Agricultural, Forestry and Food Sciences, University of Turin, Grugliasco (TO), Italy.

Correspondence:

Davide Spadaro, DISAFA, Dept. Agricultural, Forestry and Food Sciences, University of Turin, Via L. da Vinci 44, I-10095 Grugliasco (TO), Italy.

Telephone: +39-011-6708942
Fax: +39-011-6709307
E-mail: davide.spadaro@unito.it
ABSTRACT

A total of 130 spice samples coming from India, China, South America, USA, Northern Africa, Europe and Sub-Saharan Africa were collected in different stores of Northern Italy. They were analyzed for aflatoxins (AFs: AFB₁, AFB₂, AFG₁, AFG₂) and ochratoxin A (OTA) content by liquid chromatography with mass spectroscopy and positive electrospray ionization (LC/ESI-MS/MS), and HPLC with fluorescence detector (FLD), respectively. The analysis showed that 20 (15.4%) and 31 (23.8%) out of 130 samples were contaminated with AFs and OTA, respectively. A low level of total AFs contamination was found in the positive samples, the average concentration was 0.64 ng g⁻¹, far below the maximum threshold admitted by the European legislation (5 ng g⁻¹ for AFB₁, and 10 ng g⁻¹ for total aflatoxins (AFB₁, AFB₂, AFG₁ and AFG₂). A higher incidence of OTA was found in chili (60.0%) more than in pepper (13.3%), ranging from 2.16 to 16.35 ng g⁻¹, and from 1.61 to 15.85 ng g⁻¹, respectively. Moreover, three spice samples (2.3%) contaminated by OTA trespassed the threshold admitted by the European Regulation (EC, 2010). The co-occurrence of OTA and AFs in spices was detected in 6 out of 130 samples (4.6%), ranging from 1.61 to 15.85 ng g⁻¹ and from 0.57 to 3.19 ng g⁻¹, respectively.

Keywords: aflatoxins, ochratoxin A, chili, Italy, pepper.
1. Introduction

Spices are widely used all over the world for food preparation, to increase the flavour and aroma, and also in the past as food preservative. Considering the global trade value, the most cultivated spice crops are pepper, capsicum, nutmeg, cumin and cinnamon. Spices are commercially produced in a relatively low number of countries. India is the most important spice producer (74% of the world market), followed by Bangladesh (6%), Turkey (5%) and China (5%) (FAOSTAT, 2010).

The fungal growth and development on spices are favoured by environmental conditions, such as temperature and humidity, and by poor manufacturing conditions in the production region. Moulds decrease quality and quantity of food production, and may also create potential risk for human and animal health, due to the production of secondary metabolites, called mycotoxins. Some mycotoxins, like AFs and OTA, are produced by species of *Aspergillus* and *Penicillium*. These mould species can develop in post-harvest, during drying and storage, particularly when good storage practices are not adopted (Scott, 1984).

Aflatoxins (AFs, sum of aflatoxins B₁, B₂, G₁, G₂) and ochratoxin A (OTA) can be found in several types of food commodities, including spices (Wangikar, Dwivedi, Sinha, Sharma, & Telang, 2005).

AFs are difuranocoumarin compounds primarily produced by certain species of *Aspergillus*: *A. flavus*, *A. parasiticus*, *A. nomius*, and *A. tamarri* (Yabe, Nakamura, & Hamasaki, 1999). *A. flavus* and *A. parasiticus* are mainly producers of AFs: aflatoxin B₁ (AFB₁), B₂ (AFB₂), G₁ (AFG₁) and G₂ (AFG₂) (Varga, Frisvad, & Samson, 2011). Occurrence of AFs contaminations on several agricultural products, e.g. maize, wheat, rice, spices, dried fruits and hazelnuts, are worldwide reported (Grajewski, Blajet-Kosicka, Twaruzek, & Kosicki, 2012; Jackson, Kudupoje, & Yiannikouris, 2012; Prelle, Spadaro, Garibaldi, & Gullino, 2012). AFs have
been clearly identified as toxic, mutagenic, teratogenic, and carcinogenic compounds. AFB$_1$ is the most potent carcinogenic compound found in nature (Castells, Marin, Sanchis, & Ramos, 2008). The International Agency for Research of Cancer (IARC, 1993) has classified AFB$_1$ as a human carcinogen (Group I). OTA is a mycotoxin primarily produced by some strains of Aspergillus belonging to the sections Circumdati and Nigri. In particular, it is produced in warm and tropical climates by Aspergillus ochraceus, and in temperate countries by Penicillium verrucosum. (Tittlemier, Varga, Scott, & Kraska, 2011; Varga, Kevei, Rinyu, Teren, & Kozakiewicz, 1996). OTA is usually found in different foods, such as cereals, spices, coffee, wine and dried fruit (Coronel, Marin, Cano-Sancho, Ramos, & Sanchis, 2012; Spadaro, Lore, Garibaldi, & Gullino, 2010). OTA has been identified as nephrotoxic, hepatotoxic, immunotoxic and teratogenic, and classified in the group 2B by IARC (1993) as a possible carcinogen for humans.

Due to favourable conditions in tropical climates, both mycotoxins can co-contaminate some typologies of spices, not only in the field, but also during drying and storage.

The European Union fixed a maximum admitted level of 5 ng g$^{-1}$ for AFB$_1$, and 10 ng g$^{-1}$ for total AFs (AFB$_1$, AFB$_2$, AFG$_1$ and AFG$_2$) intended for chili, chili powder, paprika, white and black pepper, nutmeg, turmeric, ginger and spice mixtures containing one or more of the above-mentioned spices (European Commission, 2010). In the case of OTA, the maximum admitted level for the same spices listed before has been decreased from 30 µg kg$^{-1}$ to 15 µg kg$^{-1}$ since 1/7/2012 (European Commission, 2012).

Despite many papers reported the co-occurrence of AFs and OTA content in spices from different countries, such as Turkey (Ozbey & Kabak, 2012) and Malaysia (Jalili & Jinap, 2012), only one paper reported the AFs content in 28 spice samples marketed in Italy (Romagnoli, Menna, Gruppioni, & Bergamini, 2007).
In Italy, the consumption of spices is low (around 50%), compared to other European countries, such as France, Germany or Spain, but growing quickly, due to the recent immigration from extra-European countries and to the diffusion of ethnical restaurants (Dalpozzo, 2011).

The aim of this study was to analyse the co-occurrence of AFs and OTA in a large number of spice samples collected in different stores of Northern Italy. Two efficient and simple methods of extraction, purification and analysis were validated on several spices: AFs were simultaneous detected by liquid chromatography-tandem mass spectroscopy with electrospray ionization (LC/ESI-MS/MS), and OTA by liquid fluorescence detector (FLD).

2. Materials and methods

2.1. Chemicals and reagents

HPLC grade acetonitrile and LC-MS grade methanol, formic acid and acetic acid were purchased from Sigma-Aldrich (St Louis, MO, USA). AFs (AFB$_1$, AFB$_2$, AFG$_1$, AFG$_2$) and OTA analytical standards were purchased from Sigma-Aldrich, and dissolved in methanol and acetonitrile, respectively, to prepare working standard solutions at the concentration of 10 µg l$^{-1}$. All standard solutions were stored in the dark at 4°C. NaCl, KCl, Na$_2$HPO$_4$, KH$_2$PO$_4$, polyethylene glycol (PEG), ammonium formate, NaHCO$_3$ and Tween 20 were purchased from Merck (Merck, Darmstadt, Germany). AflaClean select and OtaClean select immunoaffinity columns were obtained from LCTech (Dorfen, Germany). HPLC eluents were degassed for 5 minutes and filtered through mixed cellulose ester 0.22 µm-filters (Advantec MFS, Pleasanton, CA, USA) before use.

2.2. Samples
A total of 130 spices coming from India, China, South America, USA, Northern Africa, Europe and Sub-Saharan area were purchased randomly in North-Western Italy stores, from September 2011 to February 2012. Due to high different typologies of spices, samples were divided in: 30 samples of pepper, 25 of hot pepper and paprika, 21 of mixed spices (such as curry and food seasoning), and 54 other spices including 15 different types of spices (cinnamon, cloves, ginger, juniper, poppy seeds, coriander, fennel, vanilla, mustard, turmeric, nutmeg, sesame, cardamom, anise, dill).

Two hundred mg of each sample were stored in plastic bags, in the dark, at low relative humidity and 4°C before the analysis. All samples, except powder spices, were pulverized using a food processor, until homogeneous. Spice sampling was done in accordance with sampling provision described on European regulation No 401/2006.

2.3. LC-MS/MS and HPLC apparatus

Liquid chromatography coupled with mass spectrometry was used to detect aflatoxins in spice samples and to confirm OTA absence in spices used for validation method. 1260 Agilent Technologies consisting of binary LC pump and a vacuum degasser; connected with a Varian autosampler Model 410 Prostar (Hansen Way, CA, USA) equipped a 100 µL loop was used as liquid chromatograph and was coupled to a triple quadrupole mass spectrometer Varian 310-MS. The analytical column used for LC separation was a Zorbax Eclipse Plus C18 (100 mm x 4.6 mm, 3.8 µm particle size, Agilent). The chromatographic conditions were: column temperature: 45°C; mobile phase consisting of eluent A (water with 20 mM ammonium formate buffer at pH 3.35) and eluent B (methanol), using a flow rate of 0.3 ml min⁻¹. A gradient elution was applied as follows: 0-5 min, 50% A / 50% B – 30% A / 70% B; 5-10 min, 30% A / 70% B – 10% A / 90% B; 10-11 min, 10% A / 90% B, and 11-15 min, 10% A / 90% B – 50% A / 50% B. Two minutes of post run was applied. The injection volume was 10
The triple quadrupole mass spectrometer was operated in the positive electrospray ionization (ESI)
mode (ESI⁺). The ionization source conditions were: needle voltage of 2.5 kV, capillary voltage of 60-77 V, source temperature of 50 °C, desolvation temperature of 350°C, cone gas flow rate of 50 psi, desolvation gas flow rate of 50 psi with nitrogen. Multiple reaction monitoring (MRM) mode of operation was used. The [M+H]⁺ ions of AFs were used as parent ions. The most intense daughter ions, resulting from collision-induced dissociation with argon, were used to detect and quantify AFs content. The argon pressure was set at 1.8 psi. The most intense daughter ions detected were: m/z 284.9 at 14 eV of collision energy (CE) for AFB₁, m/z 258.9 at 22 eV CE for AFB₂, m/z 242.9 at 18 eV CE for AFG₁, m/z 245 at 24 eV CE for AFG₂, m/z 358 at 18 eV CE and m/z 239 at 30 eV CE for OTA.

OTA was detected with a HPLC apparatus 1100 series Agilent equipped with G1311 quaternary pump, G1379 degasser, G1313A autosampler, G1316A column thermostat and G1321A FLD – Fluorescence Detector. The mobile phase consisted in an isocratic mixture of acetonitrile:water:acetic acid (49:49:2) for 15 min. Sample (30 µL) was injected into the analytical column Synergi 4u Hydro-RP (250 mm x 4.6 mm, Phenomenex) and detected using 333 and 460 nm as wavelengths for excitation and emission, respectively.

2.4. Aflatoxins extraction and clean up

In the first part of this study, concerning the validation of extraction method, two samples for each matrix, which confirmed to be aflatoxin free, were used as follows: one aliquot of the sample was analysed as such, whilst other aliquots were spiked with a known concentration of mycotoxin standard. The method for AFs extraction from hazelnut, described in a previously published paper.
(Prelle et al., 2012), was used with slight modification. For 25 g of sample, 5 g of NaCl and
125 ml of extraction solution, methanol: water (80:20) were added, and left for 2 h on a
shaker apparatus at 165 rpm. To eliminate the solid phase from the extraction solution, the
sample was filtered, first, through a Whatman No. 4 filter paper, and subsequently by using a
Whatman CA 0.45 µm syringe filter. A 10 ml aliquot of filtrate was diluted 1:4 in phosphate
buffer solution (PBS) and centrifuged at 4,000 rpm. To clean up the samples, immunoaffinity
columns (IAC) were used, by loading 10 ml of diluted sample. IAC were washed with 10 ml
of PBS solution and then with 10 ml of water and air dried. AFs were eluted with 3 ml of
methanol into an amber glass vial. The elute was evaporated at 65°C under air flow and 1 ml
of mobile phase was added to the precipitate and vortexed until dissolved. All samples were
analysed in triplicate.

2.5. Ochratoxin A extraction and clean up

The absence of OTA, avoiding positive matrix effect of fluorescence signal, was confirmed
by analysis through LC-MS/MS. The samples (4g) were put in a 50 mL centrifuge tube with
20 mL of extraction solution, methanol:water (80:20), and left for 5 minutes in ultrasonic
apparatus at 25°C. To separate the solid sample from the extraction solution, centrifugation
for 15 min at 6,000 rpm was applied. After centrifugation, the solution was diluted 1:4 with
PBS solution and filtered by cellulose acetate 0.45 µm syringe filter. Ten ml of diluted and
filtered solution were loaded into IAC and, then, 10 ml of washing solution (2.5% NaCl and
0.25% NaHCO₃) and 10 ml of ultrapure water were added into column. Before eluting with 3
ml of methanol into an amber glass vial, the column was air dried. The elute was evaporated
at 65°C under air flow and 1 ml of eluent was added to the precipitate and vortexed until
dissolved. All samples were analyzed in triplicate.
2.6. Method validation

Validation of AFs and OTA methods was optimized studying apparent recovery rate (R%), limits of detection (LOD) and quantification (LOQ), based on the IUPAC definition (Thompson, Ellison, & Wood, 2002), and, to validate the extraction method and chromatographic performances, capacity factor (k‘), repeatability of recovery (RSD) and capacity factor (RSDk‘) were calculated. Following the guidelines of Commission Decision 2002/657/EC, these parameters were validated. The recovery (R%), RSD, LOD and LOQ were determined on pepper, chili and four spices chosen by “others” (cinnamon, clove, nutmeg and sesame) and two spice by “mixed” group (curry and food seasoning). Samples were spiked with four concentrations of standard solutions of AFB$_1$ (0.5, 5, 10, 25 ng ml$^{-1}$) for pepper, cinnamon and seasoning food and (1, 5, 10, 25 ng ml$^{-1}$) for chili, clove, nutmeg, sesame and curry. Validation levels for total AFs were obtained spiking at 3, 5, 10, 25 ng ml$^{-1}$, and for OTA at 5, 10, 25, 50 ng ml$^{-1}$ with standard solutions. RSDk’ and k’ were calculated, by analysing spiked cinnamon, clove, food seasoning and pepper samples at 10 ng ml$^{-1}$ of AFB$_1$, total AFs, and OTA. Each test was performed three times.

Limits of detection (LOD) and quantification (LOQ) of each method for all mycotoxins were assessed. LOD was defined as three times the electronic baseline noise and LOQ as ten times the level of the baseline noise. The baseline noise was obtained with a blank sample for each matrix processed following the tested procedures. LOD and LOQ were calculated, analysing six blank samples and for total AFs were obtained by meaning the values of each mycotoxin. The recovery was calculated, using a protocol presented by (Matuszewski, Constanzer, & Chavez-Eng, 2003).

Capacity factors (k‘), intended as a measure of the time the sample component resides in the stationary phase relative to the time in the mobile phase, were used to assure reliability and
repeatability of chromatographic analysis and were calculated for each mycotoxin (IUPAC, 1997).

2.7. Statistical analysis

Normal distribution of toxin contents, means, standard errors and validation data were analysed with SPSS software (SPSS Institute, Inc, 2000, Version 18.0). The calibration curves used for quantification were calculated by least-squares method. Samples with a concentration of AFs and OTA higher that LOD were considered positives, whilst samples with concentrations lower than LOD were considered negatives. Mean AFs and OTA concentrations were calculated only on the positive samples higher than the LOQ. The Kruskal-Wallis test was used to compare the mean AF and OTA levels among the different typologies (pepper, chili, mixed and others), and geographical origins of spice samples, while the Mann-Whitney test was used to compare the mean OTA and AFs levels in pepper/chili samples and grain/powder of pepper and chili samples, using the null hypothesis that the levels were not different.

3. Results and discussion

This study represents the first report about monitoring of AFs and OTA co-occurrence in a large number of spice samples marketed in Italy. Two extraction and detection methods for AFs and OTA were validated on chili, pepper, cinnamon, nutmeg, food seasoning and curry matrices by LC/ESI-MS/MS and by FLD, respectively. Due to the limited consumption, and consequently limited availability of spices in Italy, with the exception of pepper and chili, we grouped the spices in four categories: pepper, chili, others and mixed. The geographical origins of the spices analysed are shown in table 1. In
agreement with a FAO report (FAOSTAT, 2012) about the main spice producers, most of the analysed samples were coming from India (74), followed by North Africa (33), and China (11).

3.1. Analytical performance

The analytical methods were validated considering: linearity, apparent recovery rate (R%), capacity factor (k’), repeatability of recovery (RSD) and capacity factor (RSDk), limits of detection (LOD) and quantification (LOQ) for AFs and OTA methods. Data of performed analytical methods are summarized in Table 2. Due to complexity of chili and pepper matrix, LOD and LOQ obtained for OTA detection, 0.86 ng g\(^{-1}\) and 2.61 ng g\(^{-1}\) for pepper, and 0.70 ng g\(^{-1}\) and 2.13 ng g\(^{-1}\) for chili, respectively, were higher than other spices tested. These results could be explained by presence of matrix compounds which interfere with analytical signal increasing baseline noise (Prelle, Spadaro, Denca, Garibaldi & Gullino, 2013). OTA recovery rate ranged from 70.8 % at 25 ng g\(^{-1}\) of sesame to 102.1 % at 10 ng g\(^{-1}\) of nutmeg. According to Castagnaro et al. (2006), we could explain these recovery results obtained as interaction between several interferences on specific identification antigen-antibody present into IAC. In particular, the specific antibodies identification could be reduced by the formation of open-ring OTA at alkaline pH and by cross-reaction due to the presence of nonchlorinated analogue OTB (Castegnaro, Tozlovanu, Wild, Molinie, Sylla, & Pfohl-Lesztkowicz, 2006). Repeatability of OTA measurements, ranging from 0.3% for curry to 18.2% for food seasoning, resulting in accordance with requirement established by EU regulation. On the contrary the values of LOD and LOQ for chili and pepper for aflatoxin detection, ranging from 0.08 ng g\(^{-1}\) of AFB1 to 0.41 of AFs, respectively, for pepper and from 0.17 ng g\(^{-1}\) of AFB1 to 0.71 of AFs, respectively, for chili resulted less influenced by matrix interferences on analytical signal than OTA detection. Recovery values for AFs and AFB1 were above of
70% for all spices tested, despite the matrix complexity of some kind of spice, such as curry and food seasoning, composed by different quantity of other spices. In the other hand the k’ and RSDk’ values obtained reported in Table 3, demonstrated the high repeatability and hardiness (RSDk’% < 0.1) of optimized chromatographic method, despite different spices analysed.

3.2. Mycotoxin occurrence

AFs were found in 20 (15.4%) out of 130 samples analysed with levels ranging from 0.59 to 5.38 ng g⁻¹. The average concentration was 0.96 ng g⁻¹. Five samples of each typology of spice (pepper, chili, mixed and others) resulted positive to one or more AFs. The AFs contamination range was between 0.59 and 3.68 ng g⁻¹, with 11 (8.5%) samples contaminated by AFB₁, with a mean on contaminated samples of 0.31 ng g⁻¹, 14 (4.6%) by AFB₂ (contamination mean: 1.07 ng g⁻¹), 3 (2.3%) by AFG₁ (0.42 ng g⁻¹), and 7 (5.4%) by AFG₂ (2.26 ng g⁻¹) (Table 4).

No sample was contaminated by AFs above the maximum admitted threshold established by the European legislation (European Commission, 2010). In fact, the highest AFB₁ level detected was 1.95 ng g⁻¹, 2.5 times lower than the maximum threshold specific for AFB₁. In contrast with the results obtained by Romagnoli et al. (2007) on monitoring of AFs on spices purchased in Italy, we detected a lower incidence of contamination and a lower level of AFs. Although pepper and chili are usually more susceptible substrates to mycotoxins contamination (Reddy, Mayi, Reddy, Thirumala-Devi, & Reddy, 2001), our results showed no differences among the spice typologies analysed. The low amount of AF content detected in our study agrees with the results reported by similar studies (Jalili & Jinap, 2012; Ozbey & Kabak, 2012). The incidence of positive samples was slightly higher in the mixed and chili samples, 23.8% and 20.0%, respectively, compared to pepper (16.6%) and other (9.26%)
spices. The Kruskal-Wallis test showed that the level of contamination of AFs in different countries were not statistically different \((p=0.586)\). Average AF content in pepper and chili samples were calculated with Mann-Whitney test, and resulted not statistically different \((p=0.248)\).

Considering the OTA contamination in the spice samples, the incidence resulted higher than AFs: 23.8% spices were positive to OTA, while only 15.4% to AFs. The values of OTA contamination ranged from LOD to 19.06 ng g\(^{-1}\). The contamination OTA mean on positive samples was 6.18 ng g\(^{-1}\). Fifteen out of 25 (60%) chili samples were contaminated at levels ranging from 2.16 to 16.35 ng g\(^{-1}\), 7 out of 21 (33.3%) mixed samples at levels ranging from 1.84 to 19.06 ng g\(^{-1}\). Pepper (13.3%) and other samples (9.3%) resulted positive ranging from 1.61 to 15.85 and from 2.21 to 11.08, respectively (Table 4).

Similarly to AFs, results of Kruskal-Wallis test showed no significant differences among the geographical origins of the samples. The Mann-Whitney test, instead, reported significant differences on OTA contamination levels between chili and pepper \((p<0.05)\). Significant differences were also noticed for grain and powder chili \((p<0.05)\). Chili powder showed higher OTA content (78.9%) than chili grains, probably, due to the higher contact surface presented to the microbial population, which can favor the mycotoxin release.

The current monitoring showed a higher incidence of OTA than AFs in spice samples marketed in Italy; moreover, concentration levels of OTA detected in three (2.3%) samples out of 130 trespassed the threshold admitted by the European Regulation (EC, 2010).

The co-occurrence of OTA and AFs in spices was detected in 6 out of 130 samples(4.6%), in particular in chili, pepper and curry samples. OTA and AFs concentration ranged from 1.61 to 15.85 ng g\(^{-1}\) and 0.57 to 3.19 ng g\(^{-1}\), respectively. Several studies were performed over the last decade on the co-occurrence of AFs and OTA in spices. Most of the studies highlighted that the reason for a high mycotoxin contamination is the susceptibility of spices to fungal
contamination, and consequently to mycotoxin production, connected to environmental
conditions, such as high humidity and temperature (Patharajan et al., 2011), of the producer
countries located in tropical and sub-tropical areas. Regarding the co-occurrence of AFs and
OTA in same substrate, Sedmikova, Reisnerova, Dufkova, Barta, & Jilek (2001)
demonstrated that OTA could increase the mutagenic activity of AFB₁. In agreement with our
results, most of previous work reported high incidence of AFs and OTA in pepper and chili.
In particular, during the last decade, AFs and OTA were monitored in over 1,500 spice
samples, resulting in a contamination incidence ranging from 30 to 70%. Despite the wide
range of spices present, attention mainly focused on pepper and chili, due to their high
demand in the market worldwide and to their susceptibility to fungal contamination,
especially during the drying processes (Shundo, de Almeida, Alaburda, Lamardo, Navas,
Ruvieri, et al., 2009). According to the available literature and to our results, chili and pepper
are more frequently contaminated by AFs and OTA than other spices, such as cumin or
cinnamon. This is the first study on AFs and OTA co-occurrence in a large number of spice
samples marketed in Italy. Future studies will also monitor the fungal population present on
the spices, which could be related to the AFs and OTA contamination, with microbiological
an molecular tools (Spadaro, Patharajan, Karthikeyan, Lorè, Garibaldi, & Gullino, 2011).

Acknowledgements

Work carried out within the framework of the project “ITACA – Technological Innovation,
Automation and new Analytical Controls to improve the quality and safety of the
Piedmontese food products” funded by the Piedmont Region (Italy).
References


FAOSTAT. (2012). Website: http://faostat.fao.org


Table 1. Geographical origin of spice samples.

<table>
<thead>
<tr>
<th>Origin</th>
<th>Pepper (n)</th>
<th>Chili (n)</th>
<th>Mixed (n)</th>
<th>Others (n)</th>
</tr>
</thead>
<tbody>
<tr>
<td>South America</td>
<td>2</td>
<td>2</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>India</td>
<td>25</td>
<td>15</td>
<td>8</td>
<td>26</td>
</tr>
<tr>
<td>North Africa</td>
<td>2</td>
<td>2</td>
<td>13</td>
<td>16</td>
</tr>
<tr>
<td>Europe</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>China</td>
<td>1</td>
<td>5</td>
<td>0</td>
<td>5</td>
</tr>
<tr>
<td>Sub-Saharan Africa</td>
<td>0</td>
<td>1</td>
<td>0</td>
<td>5</td>
</tr>
<tr>
<td>USA</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td><strong>Total samples (n)</strong></td>
<td><strong>30</strong></td>
<td><strong>25</strong></td>
<td><strong>21</strong></td>
<td><strong>54</strong></td>
</tr>
</tbody>
</table>
## Table 2. Recovery of AFs and OTA spiked into pepper and cinnamon samples.

<table>
<thead>
<tr>
<th>Analytes</th>
<th>Matrix</th>
<th>Fortification (ng ml^{-1})</th>
<th>Recovery (n=3) (%)</th>
<th>RSD (%) (n=3)</th>
<th>LOD (ng g^{-1}) (n=6)</th>
<th>LOQ (ng g^{-1}) (n=6)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Aflatoxin B_1</td>
<td>pepper</td>
<td>0.5, 5, 10, 25</td>
<td>91.8, 97.8, 99.6, 99.7</td>
<td>3.8, 0.6, 2.8, 3.5</td>
<td>0.08</td>
<td>0.28</td>
</tr>
<tr>
<td>Aflatoxins B_1, B_2, G_1, G_2</td>
<td>pepper</td>
<td>3, 5, 10, 25</td>
<td>79.1, 82.8, 82.5, 83.3</td>
<td>25.6, 23.5, 25.0, 25.4</td>
<td>0.12</td>
<td>0.41</td>
</tr>
<tr>
<td>Aflatoxin B_1</td>
<td>chili</td>
<td>1, 5, 10, 25</td>
<td>100.6, 97.3, 93.4, 90.5</td>
<td>1.8, 2.0, 2.9, 2.7</td>
<td>0.17</td>
<td>0.55</td>
</tr>
<tr>
<td>Aflatoxins B_1, B_2, G_1, G_2</td>
<td>chili</td>
<td>3, 5, 10, 25</td>
<td>78.04, 83.4, 85.5, 88.0</td>
<td>1.9, 1.1, 2.5, 1.2</td>
<td>0.21</td>
<td>0.71</td>
</tr>
<tr>
<td>Aflatoxin B_1</td>
<td>cinnamon</td>
<td>0.5, 5, 10, 25</td>
<td>99.8, 99.8, 99.6, 99.9</td>
<td>0.1, 2.7, 3.1, 3.0</td>
<td>0.12</td>
<td>0.41</td>
</tr>
<tr>
<td>Aflatoxins B_1, B_2, G_1, G_2</td>
<td>cinnamon</td>
<td>3, 5, 10, 25</td>
<td>80.5, 81.5, 81.9, 83.1</td>
<td>24.8, 25.5, 25.8, 23.7</td>
<td>0.18</td>
<td>0.61</td>
</tr>
<tr>
<td>Aflatoxin B_1</td>
<td>nutmeg</td>
<td>1, 5, 10, 25</td>
<td>79.0, 77.4, 77.6, 82.1</td>
<td>12.3, 14.3, 8.8, 8.9</td>
<td>0.2</td>
<td>0.66</td>
</tr>
<tr>
<td>Aflatoxins B_1, B_2, G_1, G_2</td>
<td>nutmeg</td>
<td>3, 5, 10, 25</td>
<td>76.9, 80.3, 89.7, 87.4</td>
<td>9.1, 5.9, 12.7, 7.3</td>
<td>0.65</td>
<td>2.2</td>
</tr>
<tr>
<td>Aflatoxin B_1</td>
<td>clove</td>
<td>1, 5, 10, 25</td>
<td>81.6, 92.7, 90.3, 85.1</td>
<td>13.3, 16.9, 12.4, 10.5</td>
<td>0.17</td>
<td>0.58</td>
</tr>
<tr>
<td>Aflatoxins B_1, B_2, G_1, G_2</td>
<td>clove</td>
<td>3, 5, 10, 25</td>
<td>83.9, 96.6, 87.3, 81.2</td>
<td>15.6, 13.0, 9.4, 11.4</td>
<td>0.57</td>
<td>1.94</td>
</tr>
<tr>
<td>Aflatoxin B_1</td>
<td>sesame</td>
<td>1, 5, 10, 25</td>
<td>73.2, 73.0, 77.8, 83.6</td>
<td>5.9, 3.2, 7.3, 5.6</td>
<td>0.19</td>
<td>0.62</td>
</tr>
<tr>
<td>Aflatoxins B_1, B_2, G_1, G_2</td>
<td>sesame</td>
<td>3, 5, 10, 25</td>
<td>82.7, 81.1, 77.3, 81.9</td>
<td>20.3, 13.2, 10.4, 12.6</td>
<td>0.33</td>
<td>1.12</td>
</tr>
<tr>
<td>Aflatoxin B_1</td>
<td>food seasoning</td>
<td>0.5, 5, 10, 25</td>
<td>97.2, 97.6, 87.8, 81.8</td>
<td>4.7, 1.1, 2.9, 1.8</td>
<td>0.13</td>
<td>0.45</td>
</tr>
<tr>
<td>Aflatoxins B_1, B_2, G_1, G_2</td>
<td>food seasoning</td>
<td>3, 5, 10, 25</td>
<td>92.8, 93.1, 94.3, 83.8</td>
<td>18.6, 6.3, 4.7, 5.1</td>
<td>0.26</td>
<td>0.88</td>
</tr>
<tr>
<td>Aflatoxin B_1</td>
<td>curry</td>
<td>1, 5, 10, 25</td>
<td>71.6, 74.7, 79.8, 80.3</td>
<td>19.6, 17.7, 15.9, 15.3</td>
<td>0.25</td>
<td>0.83</td>
</tr>
<tr>
<td>Aflatoxins B_1, B_2, G_1, G_2</td>
<td>curry</td>
<td>3, 5, 10, 25</td>
<td>73.0, 71.6, 78.6, 79.4</td>
<td>17.3, 17.1, 18.0, 17.9</td>
<td>0.77</td>
<td>2.62</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>-----</td>
<td>-----</td>
<td>-----</td>
<td>-----</td>
<td>-----</td>
<td>-----</td>
<td>-----</td>
</tr>
<tr>
<td>G$_2$</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>OTA</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>pepper</td>
<td>5, 10, 25, 50</td>
<td>91.1, 94.8, 97.8, 98.7</td>
<td>2.1, 2.9, 4.8, 2.1</td>
<td>0.86</td>
<td>2.61</td>
</tr>
<tr>
<td>OTA</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>chili</td>
<td>5, 10, 25, 50</td>
<td>81.1, 79.3, 88.4, 87.8</td>
<td>4.7, 8.6, 4.5, 6.7</td>
<td>0.70</td>
<td>2.13</td>
</tr>
<tr>
<td>OTA</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>cinnamon</td>
<td>5, 10, 25, 50</td>
<td>97.4, 96.3, 99.2, 95.6</td>
<td>3.4, 4.5, 4.5, 3.6</td>
<td>0.03</td>
<td>0.09</td>
</tr>
<tr>
<td>OTA</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>nutmeg</td>
<td>5, 10, 25, 50</td>
<td>80.2, 102.1, 84.6, 71.0</td>
<td>17.8, 7.8, 0.8, 0.4</td>
<td>0.03</td>
<td>0.10</td>
</tr>
<tr>
<td>OTA</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>clove</td>
<td>5, 10, 25, 50</td>
<td>78.4, 80.4, 75.3, 76.0</td>
<td>2.3, 17.1, 13.0, 0.68</td>
<td>0.02</td>
<td>0.07</td>
</tr>
<tr>
<td>OTA</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>sesame</td>
<td>5, 10, 25, 50</td>
<td>73.5, 74.5, 70.8, 75.4</td>
<td>6.0, 6.1, 6.9, 14.5</td>
<td>0.05</td>
<td>0.14</td>
</tr>
<tr>
<td>OTA</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>food seasoning</td>
<td>5, 10, 25, 50</td>
<td>78.1, 78.2, 82.7, 77.3</td>
<td>1.8, 14.5, 18.2, 2.5</td>
<td>0.05</td>
<td>0.15</td>
</tr>
<tr>
<td>OTA</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>curry</td>
<td>5, 10, 25, 50</td>
<td>79.6, 72.9, 78.3, 71.2</td>
<td>0.3, 8.2, 9.4, 0.8</td>
<td>0.04</td>
<td>0.12</td>
</tr>
</tbody>
</table>
Table 3. Analytical chromatographic performance.

<table>
<thead>
<tr>
<th>Analytes</th>
<th>Matrix</th>
<th>Capacity factor (k')</th>
<th>RSDk' (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>AFs</td>
<td>pepper</td>
<td>38.1, 36.6, 34.1, 31.8</td>
<td>0.06, 0.07, 0.09, 0.1</td>
</tr>
<tr>
<td>OTA</td>
<td>pepper</td>
<td>5.2</td>
<td>0.04</td>
</tr>
<tr>
<td>AFs</td>
<td>cinnamon</td>
<td>38.4, 36.7, 34.3, 31.9</td>
<td>0.05, 0.07, 0.07, 0.09</td>
</tr>
<tr>
<td>OTA</td>
<td>cinnamon</td>
<td>5.2</td>
<td>0.05</td>
</tr>
<tr>
<td>AFs</td>
<td>food seasoning</td>
<td>38.0, 36.6, 34.3, 31.7</td>
<td>0.03, 0.04, 0.03, 0.06</td>
</tr>
<tr>
<td>OTA</td>
<td>food seasoning</td>
<td>5.2</td>
<td>0.03</td>
</tr>
<tr>
<td>AFs</td>
<td>clove</td>
<td>37.9, 36.2, 34.1, 31.0</td>
<td>0.09, 0.1, 0.1, 0.12</td>
</tr>
<tr>
<td>OTA</td>
<td>clove</td>
<td>5.3</td>
<td>0.07</td>
</tr>
</tbody>
</table>
Table 4. Occurrence and levels of AFs and OTA in the spices analysed.

<table>
<thead>
<tr>
<th>Mycotoxins</th>
<th>Positive/total (%)</th>
<th>Mean of contamination ± SD (ng g⁻¹)</th>
<th>&lt; LOD</th>
<th>LOD-5.0</th>
<th>5.0-10.0</th>
<th>10.0-15.0</th>
<th>&gt;15.0</th>
</tr>
</thead>
<tbody>
<tr>
<td>AFB₁</td>
<td>11/130 (8.5%)</td>
<td>0.31±0.003</td>
<td>8</td>
<td>3</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>AFB₂</td>
<td>6/130 (4.6%)</td>
<td>1.07±0.086</td>
<td>4</td>
<td>2</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>AFG₁</td>
<td>3/130 (2.3%)</td>
<td>0.42±0.025</td>
<td>3</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>AFG₂</td>
<td>7/130 (5.4%)</td>
<td>2.26±0.191</td>
<td>3</td>
<td>4</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>OTA</td>
<td>31/130 (23.8%)</td>
<td>6.18±0.639</td>
<td>-</td>
<td>15</td>
<td>11</td>
<td>2</td>
<td>3</td>
</tr>
<tr>
<td>OTA+AFs</td>
<td>6/130 (4.6%)</td>
<td>0.85±0.030</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>