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## ADVERSE EFFECTS OF COMBINED MYCOTOXINS\*

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This article brings an overview of mycotoxin co-occurrence in foods in Croatia and neighbouring countries and experimental data from mycotoxin interaction studies involving *Fusarium* toxins, ochratoxin A (OTA), and aflatoxin B<sub>1</sub> (AFB<sub>1</sub>). Only a few studies of combined mycotoxin toxicity have employed a mathematical/statistical design, while others have used common statistics in order to compare the effects of mycotoxin mixtures with effects of single toxins. So far, most studies have observed additive or synergistic effects, suggesting that these mixtures pose a significant threat to human and animal health.

**KEY WORDS:** aflatoxin, citrinin, fumonisins, mycotoxin interactions, ochratoxin, trichothecenes, zearalenone

Mycotoxins are the secondary metabolites of moulds and invariably contaminate food and feed all over the world. Among hundreds of known mycotoxins, aflatoxins (AFs), ochratoxin A (OTA), fumonisins (FBs), zearalenone (ZEA), and trichothecenes stand out as the most common contaminants in a variety of food. Ingestion of these mycotoxins may cause acute toxicity or chronic disorders, depending on concentration and duration of exposure. Moreover, they are often responsible for financial losses in food production and livestock breeding. Most countries have legislation that prescribes maximum permissible concentrations of mycotoxins in certain types of food, but these regulations vary significantly between them. Croatia has harmonised its legislation on mycotoxins in foods with the European Union regulations (1).

Even though food is often contaminated with more than one mycotoxin, most studies are limited to the

toxicology of a single mycotoxin. This review summarises the findings on mycotoxin co-occurrence in food in Croatia and neighbouring countries and the experimental data on their combined toxicity.

### *Co-occurrence of mycotoxins in food*

Most of the earlier mycotoxin surveys were focused on the occurrence of single toxins in food. For example, studies conducted in Croatian and Bulgarian endemic nephropathy (EN) areas were focused on the levels of resident exposure to OTA (2, 3). Cereals are commonly contaminated with the *Fusarium* species and therefore often analysed for the presence of *Fusarium* mycotoxins such as FBs, ZEA, deoxynivalenol (DON), diacetoxiscitpenol (DAS), beauvericin (BEA), or T-2 toxin (4-6), but only a limited number of studies have paid attention to or specified the percentage of co-contaminated samples. Table 1. summarises the co-occurrence of various mycotoxins in food samples in Croatia and neighbouring countries over the last fifteen years.

\* The subject of this article has partly been presented at the International Symposium "Power of Fungi and Mycotoxins in Health and Disease" held in Primošten, Croatia, from 19 to 22 October 2011.

**Table 1** Co-occurrence of some mycotoxins in food from Croatia and the neighbouring countries

Sample / Year/ Country	Mycotoxin	Contamination / %	Range / µg kg <sup>-1</sup>	Co-contamination / %	Reference
Pre-harvest maize / 1992-93 / Italy*	FB <sub>1</sub> BEA	100 67	125000 to 250000 5000 to 10000	67	81
Stored maize / Hungary	FB <sub>1</sub> ZEA DON T-2	70.8 87.5 70 41.7	50 to 19800 10 to 11800 70 to 21200 60 to 390	Not specified	4
Harvested maize/ Hungary	FB <sub>1</sub> ZEA DON T-2	70 17 13 39	95 to 52400 6 to 79 50 to 118 50 to 551		
Maize / 1996-97 / Croatia**	OTA FB <sub>1</sub> BEA	29 95 11	0.26 to 614 12 to 11661 13 to 1864	Two toxins; 6 Three toxins; 2	7
Grains and feeds / 1998-2004 / Croatia***	T-2 DAS DON	16.8 27.6 41.2	100 to 700 100 to 500 100 to 3440	Not specified	6
Cereals / 1999 / Bulgaria**	OTA CTN	35 9.4	<0.5 to 140 <5 to 420	22	8
Maize / 2002 / Croatia	OTA FB <sub>1</sub> FB <sub>2</sub>	33 100 13	0.73 to 2.54 196.8 to 1377.6 68.4 to 3084	Two toxins; 55 Three toxins; 37	82
Foodstuffs / 2002 / Italy	FB <sub>1</sub> FB <sub>2</sub> DON	26 35 84	10 to 2870 10 to 790 7 to 930	Not specified	5
Spices / 2004 / Hungary	AFB <sub>1</sub> OTA	21 36	6.1 to 15.7 10.6 to 66.2	Not specified	83
Total diet Serbia**	OTA CTN	34 16	0.29 to 2.25	10	9
Maize, feed / 2007 / Croatia**	AFs OTA ZEA FB <sub>1</sub> +FB <sub>2</sub> +FB <sub>3</sub>	24.3 16.2 92 27	2.3 to 10.3 2.5 to 31.7 12.5 to 1182 200 to 20700	Two toxins; 4.2 to 54 Three toxins; 4.2 to 7.6	15
Pasta / Italy	AFB <sub>1</sub> OTA DON	Not detected 96.3 81.5	 0.2 to 0.52 35.1 to 450.0	Not specified	84

\* Only six maize samples were analysed

\*\* Samples were taken in the EN area

\*\*\* A total of 465 samples were analysed

FB<sub>1</sub> is the most frequent maize contaminant which possesses several toxic properties (Table 2). It occurs in significantly higher concentrations than any other *Fusarium* toxin. It is often accompanied in maize by ZEA, DON, or OTA. Due to its

nephrotoxicity, special attention has been paid to its co-occurrence with OTA as well as to co-occurrence of OTA and citrinin (CTN), particularly in the EN areas (2), as these areas report higher crop contamination with OTA and FB<sub>1</sub> or OTA and CTN

**Table 2** Mechanism of action and toxic properties of some mycotoxins, frequently found as food and feed contaminants.

Mycotoxins	Primary events at the cellular level	Toxic properties	References
Aflatoxin B <sub>1</sub>	Metabolic activation → AFB1-8,9-epoxide → modification of major cell macromolecules	Hepatotoxic, immunosuppressive, carcinogenic (group 1, IARC), teratogenic, mutagenic	66, 67
Beauvericin	Complex with essential cations (Ca <sup>2+</sup> , Na <sup>+</sup> , K <sup>+</sup> ) → inhibition of cation-selective channels in lipid membranes, inhibitor of cholesterol acyltransferase	Antimicrobial, insecticidal cytotoxic, ionophoric action	85
Citrinin	Affects mitochondrial permeability transition, calcium flux, and cytochrome c release from mitochondria, inhibits macromolecule biosynthesis → cell death	Nephrotoxic, hepatotoxic genotoxic (group 3, IARC), teratogenic, immunotoxic	86-88
Deoxynivalenol	Inhibition of protein synthesis → cell death, disruption of cytokine regulation	Causes nausea, food refusal, vomiting, diarrhoea, immunotoxic, IARC group 3	89-91
Diacetoxyscirpenol and T-toxin	Inhibition of protein synthesis → cell death T-2 → erythrocyte lysis, induction of lipid peroxidation, apoptosis, inhibition of mitochondrial electron transport	Alimentary toxic aleukia (inflammation of the skin, vomiting, damage to hematopoietic tissues, haemorrhagia), IARC group 3	89-92
Fumonisin B <sub>1</sub>	Inhibition of ceramide synthase → changes in sphingolipid metabolism, → protein kinase activity, oxidative stress → oxidative damage of cell macromolecules	Equine leukoencephalomalacia, porcine pulmonary oedema, hepatotoxic, nephrotoxic immunosuppressive, carcinogenic (group 2B, IARC)	89-91, 93
Ochratoxin A	Competition with phenylalanine and inhibition of Phe-dependent enzymes, inhibition of protein and DNA synthesis, mitochondrial transport system and transport of organic anions and cations, oxidative stress, DNA damage, affects glucose metabolism and Ca <sup>2+</sup> homeostasis,	Nephrotoxic, neurotoxic hepatotoxic, affects blood coagulation, immunotoxic carcinogenic (group 2B, IARC) teratogenic	9, 50, 94
Zearalenone	Resembles 17β-oestradiol → binds to oestrogen receptors in mammalian target cells	Disruption of hormonal control, IARC group 3	89, 91

than non-EN villages in Bulgaria, Croatia, and Serbia (7-9). Until recently it was thought that FBs were produced only by the *Fusarium* species that contaminate maize and that their occurrence in foods was limited to that substrate. However, black

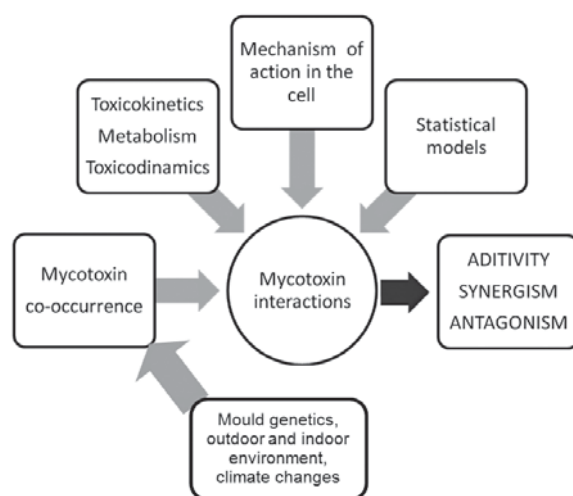
Aspergilli, including *A. niger* and *A. awamori*, which contaminate substrates with a high sugar content such as dried fruits, are able to produce FB<sub>1-4</sub>, 3-epi-FB<sub>3</sub>, 3-epi-FB<sub>4</sub>, iso-FB<sub>1</sub>, and two iso-FB<sub>2,3</sub> forms (10). Therefore, FBs and OTA could co-

contaminate these substrates as well as grapes, and both toxins could be expected in red wine (11). In addition, *A. niger* strains, which have extensively been used in biotechnology and fungal mycelium from fermentation as animal feed, can produce both toxins (12). Taking into account these investigations, FBs and OTA could be expected in various foods and not only in maize.

Mould production of mycotoxins in food is influenced by a number of factors such as temperature, water activity, substrate composition, mould physiology, or interactions with other microbes (13). All these factors are directly linked to climate change. For example, the optimal temperature for the formation of AFs is 33 °C, and these toxins are far more common in the hot tropical and subtropical regions. With current climate changes, however, AFs may soon become more common in the temperate areas of Europe as well (14). The case in point are recent reports from Croatia; AFs were found in relatively many cereal samples (24.3 %) and feed in respect to earlier studies (15, 16). The increasing incidence of AF producers (*A. flavus*) in moderate climate zones may in turn reduce the number of *Penicillium* OTA producers, as OTA production requires lower temperatures (14). The optimal temperature for the formation of *Fusarium* toxins is between 20 °C and 30 °C, and one might expect a reduction in their incidence with temperature rising above 30 °C. However, *F. verticillioides*, a producer of the FBs which frequently infect maize in Southern Europe, has been associated with dry weather during grain fill and late season rains (17). Therefore, the production of FBs will be favoured by the climate change (18). Mild temperature and rain during maize growth makes the plant susceptible to infection by *F. graminearum*, a producer of DON and ZEA. The production of ZEA is also favoured by temperature fluctuations (19). In addition, cereal infections by *Fusarium* species are spread by insects whose number is also increasing due to the climate change (14, 18). All this suggests that increased occurrence of AFs in food and co-occurrence with *Fusarium* toxins may become a food safety problem in this part of Europe in the near future.

#### *Mycotoxin interactions at the cellular level: cell models and statistics*

Combined toxicity is very hard to predict because it is influenced by several factors, including chemistry and mechanism of action, toxicodynamics and



**Figure 1** Influence of various factors on mycotoxin interactions outcome in experimental model

toxicokinetics, experimental design and endpoints of the study as well as statistical aspects (Figure 1). The best approach to studying combined toxicity is to find out how a single mycotoxin acts at the cellular level and how it interacts with another mycotoxin (20). Riley (21) has indicated where mycotoxins can interfere in the cascade of cell machinery and thus affect cellular function. For example, deregulation of calcium ions can at the same time affect mitochondrial function and activate cell signal molecules or endonucleases, which in turn can lead to the inhibition of protein synthesis or to DNA or cell membrane damage, and finally to cell death. This means that multiple mycotoxins can affect certain targets and initiate more than one event in the cell machinery leading to extremely complicated cell response (20, 21). Table 2 shows the toxic properties and toxicity mechanisms of some *Fusarium* toxins, AFs, OTA, and CTN, which frequently contaminate food in Croatia and the neighbouring countries.

Over the last few years researchers have used more and more cell cultures to study the mechanisms of mycotoxin action (cell-specific function toxicity) and to predict the outcome of mycotoxin mixtures. Cell cultures have many limitations such as immortalisation, limited survival, metabolic imbalance, or absence of tissue communication (22). Nevertheless, these *in vitro* systems are particularly useful for studying the interaction between low-molecular-weight compounds such as mycotoxins and their mixtures, and cell macromolecules. For toxicity screening, Gutleb et al. (23) recommend that several cell lines are used, as they differ in sensitivity. For example, some mammalian

cells are sensitive to fumonisin cytotoxicity. Furthermore, different cell lines can answer specific questions related to mycotoxin kinetics, active transport, metabolism, or biotransformation (22). For example, Caco-2 cell (human epithelial colorectal adenocarcinoma) layers are useful for studying absorption in the small intestine, because these cells morphologically and functionally resemble enterocytes in the small intestine (24); Hep G2 cells (human hepatocellular carcinoma) are useful for studies of liver metabolism and toxicity of xenobiotics, while artificially increased expression of specific P450 cytochromes makes them sensitive to promutagens such as aflatoxins or related mycotoxins (25). Human lung adenocarcinoma (A549) cells express phase I and phase II enzymes involved in detoxification or bioactivation of respiratory toxins (e. g. sterigmatocystin-STC and *Stachybotrys* toxins) and are morphologically and functionally consistent with type II pulmonary epithelial cells *in vivo* (26). A parallelogram approach makes it possible to compare data derived from animal cell studies and animal-derived data as well human-derived *in vitro* systems and animal cell systems (22).

Choosing the right cell model and carefully planning experimental design for studying mycotoxin interactions are essential for accurate mathematical and statistical analysis. So far, only a few mathematical/statistical models have been applied for mycotoxin mixture experiments, including the central composite design, full factorial design (27, 28), ray design (29), isobolographic analyses (30, 31), and unpaired *t*-test for the comparison between expected and measured toxicity of a mycotoxin mixture (28, 32-37). The aim of these mathematical designs is to predict combined mycotoxin effects based on the comparison between the observed and expected effects of a mycotoxin mixture. The ray design provides constant mixture ratios considering the concentration-response relationship. This design compares responses to concentration addition of individual toxins and can be related to the comparison between the expected and measured toxicity of combined mycotoxins analysed by the *t*-test. An alternative is the composite design ( $n \times n$ ), where particular mixture compositions are selected and analysed using the full factorial design to detect interactions at various mixture ratios. The isobolographic analysis is also based on the concentration-response relationship of individual and combined mycotoxins and provides a combination index as a quantitative measure of the degree of

interaction between mycotoxins in the mixture. These mathematical/statistical designs were employed for studying interactions between *Fusarium* mycotoxins (27, 30, 31), CTN, and OTA (28, 29, 32, 37), then OTA and FB<sub>1</sub> or BEA (33-36), and between AFB<sub>1</sub> and OTA or FB<sub>1</sub> (38, 39). The results of these studies will be discussed in the following sections.

#### *Interactions between Fusarium mycotoxins*

Studies of trichothecenes are mainly focused on their cytotoxic and immunomodulatory effects. Speijers and Speijers (20) have reviewed studies of interactions between trichothecenes (DON, DAS, T-2, nivalenol-NIV and moniliformin-MON) as well as between ZEA and FB<sub>1</sub>, including the one by Tajima et al. (27) and Thuvander et al. (40). In the latter study trichothecenes DON, DAS, T-2, and NIV inhibited both the proliferation of human lymphocytes and immunoglobulin (Ig) production *in vitro*. Combinations of NIV and other trichothecenes resulted in additive toxicity, while DON combined with T-2 or DAS showed a slight antagonism. Tajima et al. (27) investigated the inhibition of DNA synthesis in the mouse fibroblast cell line L-929 treated with NIV, DON, T-2, ZEA, and FB<sub>1</sub> alone and in two-toxin combinations. Single toxins showed concentration-dependent inhibitory effects in the following order: T-2>DON and NIV>ZEA, while FB<sub>1</sub> induced cell proliferation. Combinations produced additive effects at lower concentrations of individual toxins, while no additive effect was detected at high concentrations. ZEA and FB<sub>1</sub> and NIV and T-2 showed synergistic effects in the second, "screening stage" of the experiment. These particular mixtures were then separately studied using the full factorial design, which detected synergism between NIV and T-2, but did not confirm the one between ZEA and FB<sub>1</sub>. The authors concluded that the effect of mycotoxin mixture cannot be predicted solely on the basis of the effect of the individual toxins.

Several studies (41-43) found epithelial porcine kidney (PK15) cells highly sensitive to relatively low concentrations of trichothecenes and sphinganine analogues and useful in studying the toxicity of combined *Fusarium* mycotoxins (fusarochromanone-FUCH, NIV, 4-ac-NIV, 15-acNIV, scirpentriol-SCIRP, and DAS) found in rice culture extracts of *F. equiseti*. The relationship between single toxins was determined by the factor analysis based on correlations, while the relationship between the toxicity of culture extracts and contribution of toxins was studied using the

multiple regression analysis. The cytotoxicity of extracts was mainly attributed to the combination of FUCH and 15-ac-NIV, while SCIRP or DAS were also implicated in this mixture (42).

In a study by Luongo et al. (44),  $FB_1$  and  $\alpha$ -ZEA alone exerted different immunomodulatory effects on human lymphoblastoid Jurkat T cells, but in combination  $FB_1$  induced concentration-dependent proliferation while  $\alpha$ -ZEA was cytotoxic. The isobole analysis showed that this mixture potentiated  $\alpha$ -ZEA cytotoxicity. In addition, this mycotoxin combination significantly reduced interferon  $\gamma$  (INF $\gamma$ ) mRNA expression compared to  $\alpha$ -ZEA alone. Using the isobole analysis Ruiz et al. (30, 31) showed that a combination of BEA and DON or T-2, as well as the three-toxin combination had antagonistic effects in mammalian kidney epithelial Vero cells. However, these mixtures exerted both synergism (BEA+T-2 and BEA+DON+T-2) and antagonism (DON+BEA and DON+T-2) in Chinese hamster ovary CHO-K1 cells, depending on the time of exposure. Dombrink-Kurtzman (45) reported additive apoptotic effect of  $FB_1$ +BEA on turkey peripheral blood lymphocytes. In other words, the same toxins interact differently in different cells.

Only a few studies addressed combined effects of *Fusarium* toxins in animals (46, 47). A combination of DON and MON as well as of  $FB_1$  and DAS or OTA exerted additive or less than additive toxicity. Similarly, Müller et al. (48) did not observe significant interactions between low-dose  $FB_1$ , DON, and T-2 with OTA in weaning piglets. In contrast, chronic ingestion of single and combined  $FB_1$  and DON caused significant morphological and immunological changes in the intestine of piglets studied by Bracarense et al. (49). Synergy was observed in the number of goblet cells and eosinophils in the ileum, additive interactions in the expression of IL-10, TNF- $\alpha$ , and adherent proteins, less-than-additive interactions in the expression of INF- $\gamma$  and lesion scores, and antagonistic effects in goblet cells, plasma cells, eosinophils, and lymphocytes of the jejunum and in the expression of IL-1 $\beta$  and IL-6. This study suggests that ingestion of low doses of these toxins may predispose animals to infections by enteric pathogens through an interactive alteration of the intestine. The authors have based these conclusions on the statistical analysis of combined treatment versus control and single-toxin treatment, but their results have not been verified by mathematical or statistical models that compare theoretical and measured effects of mycotoxin mixtures.

#### *Mycotoxin interaction studies involving OTA*

OTA is one of the most studied mycotoxins due to its presence in a variety of food commodities (maize, wheat, beans, grapes, and wine) as well as due to its implication in the development of EN in humans (50, 51). This mycotoxin is often found in food together with FBs, ZEA, or CTN (Table 1).

OTA and CTN were studied for nephrotoxicity separately and in combination. In combination, they were dominantly synergistic both in vivo and in vitro (see review 20). When CTN or patulin (PAT) were combined with OTA or OTB in porcine kidney LLC-PK1 cells, these binary mixtures always resulted in higher toxicity that was confirmed as synergism by the full factorial analysis. The toxicity of these mixtures was not as pronounced in human proximal tubule-derived IHKE cells as in LLC-PK1 cells (28, 52). In contrast, CTN had an antagonistic effect on OTA-induced caspase 3-activation in IHKE cells, which might be explained by a decreased uptake of OTA into the cells when both toxins were applied at the same time (32). Concurrent application of OTA and CTN in PK15 cells resulted in apoptotic and necrotic synergism, while genotoxicity, determined by the micronucleus test, was either additive or antagonistic, depending on mycotoxin concentrations in the mixture. Co-treatment of PK15 cells with calcium chelator BAPTA-AM plus OTA and/or CTN showed that calcium played a significant role in both DNA damage and cell death (37). In green monkey kidney Vero cells a combination of OTA and CTN significantly increased cytotoxicity and DNA fragmentation while in mouse bone marrow cells, chromosome aberrations were observed. Compared to the effects of single toxins, the effect was synergistic (53). Genotoxic synergism was also observed in rats fed with OTA and CTN and in human kidney (HK2) cells (54, 55). In the kidneys of animals receiving both toxins, the amount of DNA-OTA adducts (dG-OTA) increased ten times. In HK2 cells it increased two times. Co-exposure to CTN and OTA significantly increased cyclooxygenase (COX2) and lipooxygenase (LOX) expression, which corresponded to the increase in dG-OTA adduct formation. This study supports previous findings that OTA-mediated DNA adducts are under the control of biotransformation enzymes such as CYP450 1B1, 2C9, COX, and LOX (9, 50). Before these studies, it was believed that OTA poorly metabolised and did

not form reactive intermediates capable of interacting with DNA (50). However, mass spectrometry studies have shown that OTA-derived quinone/hydroquinone redox couple is involved in the generation of DNA adducts (56, 57). This genotoxic synergism might play a role in the development of EN and related chronic kidney diseases and carcinomas.

Beside the nephrotoxic potential, OTA in combination with CTN, PAT, and gliotoxin was also studied for immunotoxic effects in human peripheral blood mononuclear cells (29). The ray design showed that lower concentrations of these toxins in mixtures caused stronger inhibition of interferon- $\gamma$  release from T-cells and natural killer cells.

The last decade has seen intensive research of the combined effects of OTA and FB<sub>1</sub> (33-36, 58-65). In most cases, these toxins interacted in a synergistic or at least additive manner. Cytotoxic synergism was observed in rat brain glioma C6 cells, human intestinal Caco-2 cells, and Vero cells. These cells were exposed to low FB<sub>1</sub> and high OTA concentrations, which showed cytotoxicity above additive effects in respect to the individual toxins (58). Subcytotoxic concentrations of OTA and FB<sub>1</sub> or OTA and BEA in the mixture additively increased lipid peroxidation and decreased the level of glutathione in PK15 cells. At the same time, these mixtures synergistically induced caspase-3 (33, 34). All three mycotoxins applied at subcytotoxic concentrations induced DNA damage in PK15 cells (measured by the micronucleus and/or comet assay), and their combinations mostly produced an additive genotoxic effect (35, 36). The ionophoric activity of BEA is probably responsible for changes in membrane permeability, facilitating the penetration of FB<sub>1</sub> and OTA into the cell. In a study by Domijan et al. (59) both the Fpg-modified and standard alkaline comet assay showed that OTA+FB<sub>1</sub> synergistically induced DNA damage in the kidneys of male Wistar rats, indicating that beside oxidative stress some other mechanism is also involved in their genotoxic effect. A recent study by Hadjeba-Medjdoub et al. (60) has shown that FB<sub>1</sub> promotes OTA-specific DNA adducts, including C-C8dG OTA adduct and OTHQ-related adduct. These specific adducts are found in human urothelial tumours in the EN areas.

In pig studies (61-63), a mixture of OTA and FB<sub>1</sub> caused stronger lesions in the kidneys, more pronounced changes in biochemical parameters, and disturbances in humoral immune response in doses that corresponded to those found in cases of porcine

nephropathy in Bulgaria and South Africa. These findings suggested that mixtures of OTA and FB<sub>1</sub> were involved in the aetiology of the disease.

Stoev et al. (64) found that beside FB<sub>1</sub> and CTN, penicillic acid (PA) also co-occurred with OTA in swine feed. PA inhibited carboxypeptidase, the enzyme involved in detoxification of OTA, which may have enhanced OTA toxicity. This and another study by the same authors (65) showed that a combination of OTA+PA produced synergistic toxicity, expressed as degenerative changes of internal organs in chickens and pigs. Experimental administration of these toxins provoked degeneration that corresponded to porcine nephropathy, which suggested that the disease may be owed to a combined action of two or more mycotoxins.

#### *Mycotoxin interaction studies involving aflatoxins*

Toxic properties of AFs have well been documented over the past five decades in both *in vivo* and *in vitro* models (66-69). Of the major AFs (AFB<sub>1</sub>, AFB<sub>2</sub>, AFG<sub>1</sub>, AFG<sub>2</sub>), AFB<sub>1</sub> is the most prevalent and most toxic. Its toxicity and carcinogenicity are linked to the metabolic conversion by the cytochrome p450 to a highly reactive AFB<sub>1</sub>-8,9-epoxide, which binds to the DNA, RNA, and proteins. AFB<sub>1</sub> co-occurred with FB<sub>1</sub> in a high-incidence area of human primary hepatocellular carcinoma in Haimen and Guangxi (Republic of China), suggesting that the mixture may be involved in the development of the disease (68, 69). This is why this combination has received the greatest attention over the last decade. Studies on growing barrows and turkey poults fed with this binary mixture resulted in additive to synergistic effects in the barrows and less-than-additive to additive effects in the poults as compared to each mycotoxin alone (70, 71). Pozzi et al. (72) reported a synergistic effect on the reduction of body weight gain in male Wistar rats receiving oral AFB<sub>1</sub> and FB<sub>1</sub>. Galderblom et al. (73) found that in the liver of male Fischer rats, FB<sub>1</sub> exerted between 200 and 400 times lower cancer initiating potential than AFB<sub>1</sub>. However, administration of FB<sub>1</sub> three weeks after AFB<sub>1</sub> resulted in a synergistic increase in hepatocyte nodules and foci. In contrast, Carlson et al. (74) reported no cancer-initiating properties of FB<sub>1</sub> in rainbow trout but enhanced tumour formation, which points to its cancer-promoting activity. Gelderblom et al. (75) have suggested that the inhibition of cell proliferation is an important mechanism for the cancer-promoting activity of FB<sub>1</sub>. These studies showed that AFB<sub>1</sub> and FB<sub>1</sub> interacted



synergistically in both cancer initiation and promotion, depending on intake conditions.

In a study by Theumer et al. (76), subchronic doses of the AFB<sub>1</sub>+FB<sub>1</sub> mixture induced more pronounced apoptosis in the liver of male Wistar rats than either toxin alone. In addition, this combination provoked a significant increase in Sa and Sa/So ratio in the kidney and liver compared to FB<sub>1</sub> alone, indicating that AFB<sub>1</sub> enhanced FB<sub>1</sub>-induced impairments of sphingolipid metabolism. In another study by Theumer et al. (77) the toxins, applied alone or in combination, affected the immune response of rats differently. Spleen mononuclear cells (SMC) of rats fed with AFB<sub>1</sub>+FB<sub>1</sub> produced higher levels of IL-4 and lower levels of IL-10 compared to the SMC of animals fed with AFB<sub>1</sub> alone. This suggests that subchronic doses AFB<sub>1</sub> and FB<sub>1</sub> may produce an imbalance in TH<sub>1</sub>-TH<sub>2</sub> cellular subpopulations. Furthermore, peritoneal macrophages in rats exposed to AFB<sub>1</sub> alone released less H<sub>2</sub>O<sub>2</sub> than in those exposed to a toxin mixture. The mechanism of these actions is still unknown. In yet another study, Theumer et al. (78) used the alkaline comet assay and the micronucleus test to measure DNA damage in SMC in Wistar rats exposed to subchronic doses of AFB<sub>1</sub>, FB<sub>1</sub>, and their mixture. All three induced a significant damage. The parameters of oxidative stress (malondialdehyde, catalase, and superoxide dismutase) were also elevated, supporting the assumption that FB<sub>1</sub> causes oxidative stress. DNA damage was more pronounced in animals fed with AFB<sub>1</sub> or a mixture of AFB<sub>1</sub> and FB<sub>1</sub> than in rats exposed to FB<sub>1</sub> alone, indicating that the DNA lesions were produced mainly by AFB<sub>1</sub>.

In a study by McKean et al. (39), a combination of AFB<sub>1</sub> at the  $\frac{3}{4}$  LD<sub>50</sub> and FB<sub>1</sub> at a dose that did not provoke acute toxicity resulted in 100 % mortality in F-344 rats, which could be considered a synergistic effect. Using an interactive index the authors confirmed that the toxic effect was at the margin of synergism. Mosquitofish (*Gambusia affinis*) and human bronchial epithelial BEAS-2B cells showed similar findings, while in HepG2 cells this binary mixture produced antagonistic effect. In another study (79), McKean et al. used the same experimental and mathematical model to establish the combined toxicity of AFB<sub>1</sub> and T-2. The two toxins interacted mostly in an additive manner, while a synergistic interaction was noted in BEAS-2B cells.

In chicks fed with a mixture of AFB<sub>1</sub> and OTA, Huff et al. (38) established a synergistic effect on growth inhibition and mortality. The primary effect of

this interaction was nephrotoxicity but not hepatotoxicity. Recently, this combination was tested in cultured monkey kidney Vero cells (94). It significantly decreased cell viability, increased DNA fragmentation and p53 activation, and decreased the expression of the anti-apoptotic factor bcl-2. The interaction between these toxins was additive.

## CONCLUSIONS

Through food humans and animals are constantly exposed to combinations of mycotoxins in a variety of concentrations. Current climate changes have influenced the distribution of mycotoxin producers and consequently the presence of particular mycotoxins as food contaminants (e.g. AFs). Exposure to high mycotoxin concentrations often provokes acute symptoms, which are seldom recognised as mycotoxicoses. Chronic intake of sub-toxic mycotoxin concentrations might alter various events at the cellular level, leading to biochemical and immunological impairments and possibly cancer. The relationship between chronic conditions and exposure to mycotoxins is even more difficult to recognise. In general, most of the mycotoxin mixture studies have observed additive or synergistic interactions, calling our attention to the significant threat to human and animal health. One of the preventive measures should be to revise current legislation on exposure limits, taking into account these new data. Future research may expand to include other relevant mycotoxins or combinations.

## REFERENCES

1. Pravilnik o najvećim dopuštenim količinama određenih kontaminanata u hrani [Regulations on maximum levels for certain contaminants in foodstuffs, in Croatian]. Narodne novine 154/2008.
2. Peraica M, Domijan A-M, Miletić-Medved M, Fuchs R. The involvement of mycotoxins in the development of endemic nephropathy. Wien Klin Wochenschr 2008;120:402-7.
3. Pepeljnjak S, Šegvić Klarić M. "Suspects" in etiology of endemic nephropathy: Aristolochic acid *versus* mycotoxins. Toxins 2010;2:1414-27.
4. Fazekas B, Kis M, Hajdu ET. Data on the contamination of maize with fumonisin B<sub>1</sub> and other fusariotoxins in Hungary. Acta Vet Hung 1996;44:25-37.
5. Cirillo T, Ritieni A, Galvano F, Amodio Cocchieri R. Natural co-occurrence of deoxynivalenol and fumonisins B<sub>1</sub> and B<sub>2</sub> in Italian marketed foodstuffs. Food Addit Contam 2003;20:566-71.

6. Sokolović M, Šimpraga B. Survey of trichothecene mycotoxins in grains and animal feed in Croatia by thin layer chromatography. *Food Contr* 2006;17:733-40.
7. Jurjević Ž, Solfrizzo M, Cvjetković B, De Girolamo A, Visconti A. Occurrence of beauvericin in corn from Croatia. *Food Technol Biotechnol* 2002;40:91-4.
8. Vrabcheva T, Usleber E, Dietrich R, Märtlbauer E. Co-occurrence of ochratoxin A and citrinin in cereals from bulgarian villages with a history of Balkan endemic nephropathy. *J Agric Food Chem* 2000;48:2483-8.
9. Pfohl-Leszkowicz A, Tozlovanu M, Manderville R, Peraica M, Castegnaro M, Stefanovic V. New molecular and field evidences for the implication of mycotoxins but not aristolochic acid in human nephropathy and urinary tract tumor. *Mol Nutr Food Res* 2007;51:1131-46.
10. Varga J, Kocsubé S, Suri S, Szigeti GY, Szekeres A, Varga M, Tóth B, Bartók T. Fumonisin contamination and fumonisin producing black *Aspergilli* in dried vine fruits of different origin. *Int J Food Microbiol* 2010;143:143-9.
11. Logrieco A, Ferracane R, Visconti A, Ritieni A. Natural occurrence of fumonisin B<sub>2</sub> in red wine from Italy. *Food Addit Contam Part A Chem Anal Control Expo Risk Assess* 2010;27:1136-41.
12. Frisvad JC, Larsen TO, Thrane U, Meijer M, Varga J, Samson RA, Nielsen KF. Fumonisin and ochratoxin production in industrial *Aspergillus niger* strains. *Plos One* 2011;6:1-6.
13. Garcia D, Ramos AJ, Sanchis V, Marin S. Predicting mycotoxins in foods: A review. *Food Microbiol* 2009;26:757-69.
14. Paterson RRM, Lima N. How will climate change affect mycotoxins in food? *Food Res Int* 2010;43:1902-14.
15. Šegvić Klarić M, Cvetnić Z, Pepeljnjak S, Kosalec I. Co-occurrence of aflatoxins, ochratoxin A, fumonisins, and zearalenone in cereals and feed, determined by competitive direct enzyme-linked immunosorbent assay and thin-layer chromatography. *Arh Hig Rada Toksikol* 2009;60:427-34.
16. Pepeljnjak S, Šegvić M. An overview of mycotoxins and toxigenic fungi in Croatia. In: Logrieco A, Visconti A, editors. *An overview on toxigenic fungi and mycotoxins in Europe*. Dordrecht: Kluwer Academic Publishers; 2004. p. 33-50.
17. Munkvold GP, Desjardins AE. *Fusarium* in maize: can we reduce their occurrence? *Plant Dis* 1997;81:556-65.
18. Miraglia M, Marvin HJP, Kleter GA, Battilani P, Brera C, Coni E, Cubadda F, Croci L, De Santis B, Dekkers S, Filippi L, Hutjes RWA, Noordam MY, Pisante M, Piva G, Prandini A, Toti L, van den Born GJ, Vespermann A. Climate change and food safety: An emerging issue with special focus on Europe. *Food Chem Toxicol* 2009;47:1009-21.
19. Dojin R, Bullerman LB. Effect of cycling temperatures on the production of deoxynivalenol and zearalenone by *Fusarium graminearum* NRRL 5883. *J Food Prot* 1999;62:1451-5.
20. Speijers GJA, Speijers MHM. Combined toxic effect of mycotoxins. *Toxicol Lett* 2004;153:91-8.
21. Riley RT. Mechanistic interactions of mycotoxins: theoretical consideration. In: Sinha, KK, Bhatanagar D, editors. *Mycotoxins in agriculture and food safety*. Basel, New York: Marcel Dekker Inc; 1998. p. 227-54.
22. Eisenbrand G, Pool-Zobel B, Baker V, Balls M, Blaauboer BJ, Boobis A, Carere A, Kevekordes S, Lhuguenot JC, Pieters R, Kleiner J. Methods of *in vitro* toxicology. *Food Chem Toxicol* 2002;40:193-236.
23. Gutleb AC, Morrison E, Murk AJ. Cytotoxicity assay for mycotoxins produced by *Fusarium* strains: a review. *Environ Toxicol Pharmacol* 2002;11:309-20.
24. Sambuy Y, De Angelis I, Ranaldi G, Scarino ML, Stamatii A, Zucco F. The Caco-2 cell line as a model of the intestinal barrier: influence of cell and culture-related factors on Caco-2 cell functional characteristics. *Cell Biol Toxicol* 2005;21:1-26.
25. Wilkening S, Stahl F, Bader A. Comparison of primary human hepatocytes and hepatoma cell line hepg2 with regard to their biotransformation properties. *Drug Metab Dispos* 2003;31:1035-42.
26. Castell JV, Donato MT, Gómez-Lechón MJ. Metabolism and bioactivation of toxicants in the lung. The *in vitro* cellular approach. *Exp Toxicol Pathol* 2005;57:189-204.
27. Tajima O, Schoen ED, Feron VJ, Groten JP. Statistically designed experiments in a tiered approach to screenmixtures of *Fusarium* mycotoxins for possible interactions. *Food Chem Toxicol* 2002;40:685-95.
28. Heussner AH, Dietrich DR, O'Brien E. *In vitro* investigation of individual and combined cytotoxic effects of ochratoxin A and other selected mycotoxins on renal cells. *Toxicol In Vitro* 2006;20:332-41.
29. Tammer B, Lehmann I, Nieber K and Altenburger R. Combined effects of mycotoxin mixtures on human T cell function. *Toxicol Lett* 2007;170:124-33.
30. Ruiz M-J, Macáková P, Juan-García A, Font G. Cytotoxic effects of mycotoxin combinations in mammalian kidney cells. *Food Chem Toxicol* 2011;49:2718-24.
31. Ruiz M-J, Franzova P, Juan-García A, Font G. Toxicological interactions between the mycotoxins beauvericin, deoxynivalenol and T-2 toxin in CHO-K1 cells *in vitro*. *Toxicol* 2011;58:315-26.
32. Knecht A, Schwerdt G, Gekle M, Humpf H-U. Combinatory effects of citrinin and ochratoxin A in immortalized human proximal tubule cells. *Mycotox Res* 2005;21:176-81.
33. Šegvić Klarić M, Pepeljnjak S, Domijan A-M, Petrik J. Lipid peroxidation and glutathione levels in porcine kidney PK15 cells after individual and combined treatment with fumonisin B<sub>1</sub>, beauvericin and ochratoxin A. *Basic Clin Pharmacol Toxicol* 2007;100:157-64.
34. Šegvić Klarić M, Rumora L, Ljubanović D, Pepeljnjak S. Cytotoxicity and apoptosis induced by fumonisin B<sub>1</sub>, beauvericin and ochratoxin A in porcine kidney PK15 cells: effects of individual and combined treatment. *Arch Toxicol* 2008;82:247-55.
35. Šegvić Klarić M, Pepeljnjak S, Rozgaj R. Genotoxicity of fumonisin B<sub>1</sub>, beauvericin and ochratoxin A in porcine kidney PK15 cells: effects of individual and combined treatment. *Croat Chem Acta* 2008;81:139-46.
36. Šegvić Klarić M, Daraboš D, Rozgaj R, Kašuba V, Pepeljnjak S. Beauvericin and ochratoxin A genotoxicity evaluated using the alkaline comet assay: single and combined genotoxic action. *Arch Toxicol* 2010;84:641-50.
37. Šegvić Klarić M, Želježić D, Rumora L, Peraica M, Pepeljnjak S, Domijan A-M. A potential role of calcium in apoptosis and aberrant chromatin forms in porcine kidney PK15 cells induced by individual and combined ochratoxin A and citrinin. *Arch Toxicol* 2012;86:97-107.
38. Huff WE, Doerr JA. Synergism between aflatoxin and ochratoxin A in broiler chickens. *Poult Sci* 1981;60:550-5.

39. McKean C, Tang L, Tang M, Billam M, Wang Z, Theodorakis CW, Kendall RJ, Wang J-S. Comparative acute and combinative toxicity of aflatoxin B<sub>1</sub> and fumonisin B<sub>1</sub> in animals and human cells. *Food Chem Toxicol* 2006;44:868-76.
40. Thuvander A, Wikman C, Gadhasson I. *In vitro* exposure of human lymphocytes to trichothecenes: Individual variation on sensitivity and effects of combined exposure on lymphocyte function. *Food Chem Toxicol* 1999;37:639-48.
41. Yike I, Allan T, Sorenson WG, Deraborn DG. Highly sensitive protein translation assay for trichothecene toxicity in airborne particulates: comparison with cytotoxicity assays. *Appl Environ Microbiol* 1999;65:88-94.
42. Morrison E, Rundberget T, Kosiak B, Aastveit Ah, Bernhoft A. Cytotoxicity of trichothecenes and fusarochromanone produced by *Fusarium equiseti* strains isolated from Norwegian cereals. *Mycopathologia* 2002;153:49-56.
43. Uhlig S, Gutleb AC, Thrane U, Flåøyen A. Identification of cytotoxic principles from *Fusarium avenaceum* using bioassay-guided fractionation. *Toxicol* 2005;46:150-9.
44. Luongo D, Severino L, Bergamo P, De Luna R, Lucisano A, Rossi M. Interactive effects of fumonisin B<sub>1</sub> and a-zearalenol on proliferation and cytokine expression in Jurkat T cells. *Toxicol In Vitro* 2006;20:1403-10.
45. Dombink-Kurtzman MA. Fumonisin and beauvericin induce apoptosis in turkey peripheral blood lymphocytes. *Mycopathologia* 2003;156:357-64.
46. Morris CM, Li YC, Ledoux DR, Bermudez AJ, Rottinghaus GE. The individual and combined effects of feeding moniliformin, supplied by *Fusarium fujikuroi* culture material, and deoxynivalenol in young turkey poults. *Poult Sci* 1999;78:1110-5.
47. Kubena LF, Edrington TS, Harvey RB, Phillips TD, Sarr AB, Rottinghaus GE. Individual and combined effects of fumonisin B<sub>1</sub> present in *Fusarium moniliforme* culture material and diacetoxyscirpenol or ochratoxin A in turkey poults. *Poult Sci* 1997;76:256-64.
48. Müller G, Kielstein P, Rosner H, Berndt A, Heller M, Köhler H. Studies on the influence of combined administration of ochratoxin A, fumonisin B<sub>1</sub>, deoxynivalenol and T-2 toxin on immune and defence reactions in weaner pigs. *Mycoses* 1999;42:485-93.
49. Bracarense A-PFL, Luciola J, Grenier B, Pacheco GD, Moll W-D, Schatzmayr G, Oswald IP. Chronic ingestion of deoxynivalenol and fumonisin, alone or in interaction, induces morphological and immunological changes in the intestine of piglets. *Br J Nutr* 2012;107:1776-86.
50. Pfohl-Leskowicz A, Manderville RA. Ochratoxin A: an overview on toxicity and carcinogenicity in animals and humans. *Mol Nutr Food Res* 2007;51:61-99.
51. Peraica M, Flajs D, Domijan A-M, Ivić D, Cvjetković B. Ochratoxin A contamination of food from Croatia. *Toxins* 2010;2:2098-105.
52. Heussner AH, O'Brien E, Haehnlein J, Biester MA, Dietrich DR. Comparison of interactive cytotoxic effects of selected mycotoxins on renal cells. *Toxicol Sci* 2004;78(Suppl 1):89.
53. Bouslimi A, Bouaziz C, Ayed-Boussema I, Hassen W, Bacha H. Individual and combined effects of ochratoxin A and citrinin on viability and DNA fragmentation in cultured Vero cells and chromosome aberrations in mice bone marrow cells. *Toxicology* 2008;251:1-7.
54. Manderville RA, Pfohl-Leskowicz A. Bioactivation and DNA adduction as a rationale for ochratoxin A carcinogenesis. *World Mycotoxin J* 2008;1:357-67.
55. Pfohl-Leskowicz A, Molinić A, Tozlovanu M, Manderville RA. Combined toxic effects of ochratoxin A and citrinin, *in vitro* and *in vivo*. In: Siantar DP, Trucksess MW, Scott PM, Herman EM, editors. Food contaminants, mycotoxins and food allergen. Washington (D)C; American Chemical Society; 2008. p. 56-80.
56. Tozlovanu M, Faucet-Marquis V, Pfohl-Leskowicz A, Manderville RA. Genotoxicity of the hydroquinone metabolite of ochratoxin A: Structure-activity relationships for covalent DNA adduction. *Chem Res Toxicol* 2006;19:1241-7.
57. Mantle P, Faucet-Marquis V, Manderville R, Sciqualli B, Pfohl-Leskowicz A. Structures of covalent adducts between DNA and ochratoxin A: a new factor in debate about genotoxicity and human risk assessment. *Chem Res Toxicol* 2010;23:89-98.
58. Creppy EE, Chirappa P, Baudrimont I, Borracci P, Moukha S, Carratù MR. Synergistic effects of fumonisin B<sub>1</sub> and ochratoxin A: are *in vitro* cytotoxicity data predictive of *in vivo* acute toxicity? *Toxicology* 2004;201:115-23.
59. Domijan A-M, Želježić D, Kopjar N, Peraica M. Standard and Fpg-modified comet assay in kidney cells of ochratoxin A- and fumonisin B<sub>1</sub>-treated rats. *Toxicology* 2006;222:53-9.
60. Hadjeba-Medjdoub K, Faucet-Marquis V, Tozlovanu M, Peraica M, Manderville RA, Pfohl-Leskowicz A. Synergistic effect of three nephrotoxic and carcinogenic mycotoxins (citrinin, fumonisin, ochratoxin A) on human kidney cells viability and genotoxicity. In: Antolović R, Miličević T, editors. Symposium "Power of fungi and mycotoxins in health and disease"; 19-22 Oct 2011; Primošten, Croatia 2011. Book of abstracts p. 57.
61. Stoev SD, Gundasheva D, Zarkov I, Mircheva T, Zapryanova D, Denev S, Mitev Y, Daskalov H, Dutton M, Mwanza M, Schneider Y-J. Experimental mycotoxic nephropathy in pigs provoked by a mouldy diet containing ochratoxin A and fumonisin B<sub>1</sub>. *Exp Toxicol Pathol* 2012;64:733-41.
62. Stoev SD, Dutton M, Njobeh P, Mosonik J, Steenkamp P. Mycotoxic nephropathy in Bulgarian pigs and chickens: complex aetiology and similarity to Balkan Endemic Nephropathy. *Food Addit Contam A Chem Anal Control Expo Risk Assess.* 2010;27:72-88.
63. Stoev SD, Denev S, Dutton M, Njobeh P, Mosonik J, Steenkamp P, Petkov I. Complex etiology and pathology of mycotoxic nephropathy in South African pigs. *Mycotox Res* 2010;26:31-46.
64. Stoev SD, Vitanov S, Angelov G, Petkova-Bocharova T, Creppy EE. Experimental mycotoxic nephropathy in pigs provoked by a mouldy diet containing ochratoxin A and penicillic acid. *Vet Res Commun* 2001;25:205-23.
65. Stoev SD, Stefanov M, Denev S, Radić B, Domijan A-M, Peraica M. Experimental mycotoxicosis in chickens induced by ochratoxin A and penicillic acid and intervention by natural plant extracts. *Vet Res Commun* 2004;28:727-46.
66. Riley RT, Norred WP. Mechanisms of mycotoxicity. In: Howard DH, Miller JD, editors. The mycota. Vol VI. Berlin: Springer; 1996: p. 194-5.
67. International Agency for Research on Cancer (IARC). Some traditional herbal medicines, some mycotoxins, naphthalene

- and styrene. IARC Monographs on the Evaluation of Carcinogenic Risks to Humans. Vol. 82. Lyon: IARC Press; 2002.
68. Ueno Y, Iijima K, Wang S-D, Suguira Y, Sekijima M, Tanaka T, Chen C, Yu S-Z. Fumonisin as a possible contributing risk factor for primary liver cancer: a 3 year study of corn harvested in Haimen, China by HPLC and ELISA. *Food Chem Toxicol* 1997;35:1143-50.
  69. Li FQ, Yoshizawa T, Kawamura O, Luo XY, Li YW. Aflatoxins and fumonisins in corn from the high-incidence area for human hepatocellular carcinoma in Guangxi, China. *J Agric Food Chem* 2001;49:4122-26.
  70. Harvey RB, Edrington TS, Kubena LF. Influence of aflatoxin and fumonisin B<sub>1</sub> containing culture material on growing barrows. *Am J Vet Res* 1995;56:1668-72.
  71. Kubena LF, Edrington TS, Kampsholtzaple C, Harvey RB, Elissalde MH, Rottinghaus GE. Effects of feeding fumonisin B<sub>1</sub> present in *Fusarium moniliforme* culture material and aflatoxin singly and in combination to Turkey poults. *Poultry Sci* 1995;74:1295-303.
  72. Pozzi CR, Correa B, Xavier JG, Direito GM, Orsi RB, Matarazzo SV. Effects of prolonged oral administration of fumonisin B<sub>1</sub> and aflatoxin B<sub>1</sub> in rats. *Mycopathologia* 2001;151:21-7.
  73. Gelderblom WCA, Marasas WFO, Lebepe-Mazur S, Swanevelder S, Vessey CJ, Hall P de la M. Interaction of fumonisin B<sub>1</sub> and aflatoxin B<sub>1</sub> in a short term carcinogenesis model in rat liver. *Toxicology* 2002;171:161-73.
  74. Carlson DB, Williams DE, Spitsbergen JM, Ross PF, Bacon CW, Meredith FI, Riley RT. Fumonisin B<sub>1</sub> promotes aflatoxin B<sub>1</sub> and *N*-methyl-*N* Nitro-nitrosoguanidine initiated liver tumors in rainbow trout. *Toxicol Appl Pharmacol* 2001;172:29-36.
  75. Gelderblom WCA, Abel S, Smuts CM, Marnewick J, Marasas WFO, Lemmer EW, Ramljak D. Fumonisin-induced hepatocarcinogenesis: mechanisms related to cancer initiation and promotion. *Environ Health Perspect* 2001;109(Suppl 2):291-300.
  76. Theumer MG, López AG, Aoki MP, Cánepa MC, Rubinstein HR. Subchronic mycotoxicoses in rats. Histopathological changes and modulation of the sphinganine to sphingosine (Sa/So) ratio imbalance induced by *Fusarium verticillioides* culture material, due to the coexistence of aflatoxin B<sub>1</sub> in the diet. *Food Chem Toxicol* 2008;46:967-77.
  77. Theumer MG, López AG, Masih DT, Chulze SN, Rubinstein HR. Immunobiological effects of AFB<sub>1</sub> and AFB<sub>1</sub>-FB<sub>1</sub> mixture in experimental subchronic mycotoxicoses in rats. *Toxicology* 2003;186:159-70.
  78. Theumer MG, Cánepa MC, López AG, Mary VS, Dambolena JS, Rubinstein HR. Subchronic mycotoxicoses in Wistar rats: Assessment of the *in vivo* and *in vitro* genotoxicity induced by fumonisins and aflatoxin B<sub>1</sub>, and oxidative stress biomarkers status. *Toxicology* 2010;268:104-10.
  79. McKean C, Tang L, Billam M, Tang M, Theodorakis CW, Kendall RJ, Wang JS. Comparative acute and combinative toxicity of aflatoxin B<sub>1</sub> and T-2 toxin in animals and immortalized human cell lines. *J Appl Toxicol* 2006;26:139-47.
  80. Golli-Bennour EE, Kouidhi B, Bouslimi A, Abid-Essefi S, Hassen W, Bacha H. Cytotoxicity and genotoxicity induced by aflatoxin B<sub>1</sub>, ochratoxin A, and their combination in cultured Vero cells. *J Biochem Mol Toxicol* 2010;24:42-50.
  81. Bottalico A, Logrieco A, Ritieni A, Moretti A, Randazzo G, Corda P. Beauvericin and fumonisin B<sub>1</sub> in preharvest *Fusarium moniliforme* maize ear rot in Sardinia. *Food Addit Contam* 1995;12:599-607.
  82. Domijan A-M, Peraica M, Jurjević Ž, Ivić D, Cvjetković B. Fumonisin B<sub>1</sub>, fumonisin B<sub>2</sub>, zearalenone and ochratoxin A contamination of maize in Croatia. *Food Addit Contam* 2005;22:677-80.
  83. Fazekas B, Tar A, Kovács M. Aflatoxin and ochratoxin A content of spices in Hungary. *Food Addit Contam* 2005;22:856-63.
  84. Raiola A, Meca G, Mañes J, Ritieni A. Bioaccessibility of deoxynivalenol and its natural co-occurrence with ochratoxin A and aflatoxin B<sub>1</sub> in Italian commercial pasta. *Food Addit Contam Part A Chem Anal Control Expo Risk Assess* 2012;50:280-7.
  85. Šegvić Klarić M, Pepeljnjak S. Bovericin: kemizam, biološki aspekti i raširenost. [Beauvericin: chemical and biological aspects and occurrence, in Croatian]. *Arh Hig Rada Toksikol* 2005;56:343-50.
  86. Chagas GM, Campello AP, Klüppel ML. Mechanism of citrinin-induced dysfunction of mitochondria. I. Effects on respiration enzyme activities and membrane potential of renal cortical mitochondria. *J Appl Toxicol* 1992;12:123-9.
  87. Da Lozo EJ, Oliveira MB, Carnieri EG. Citrinin-induced mitochondrial permeability transition. *J Biochem Mol Toxicol* 1998;12:291-7.
  88. International Agency for Research on Cancer (IARC). Some naturally occurring substances: food items and constituents, heterocyclic aromatic amines and mycotoxins. IARC Monographs on the Evaluation of Carcinogenic Risks to Humans. Lyon: IARC; 1993.
  89. Yazar S, Omurtag GZ. Fumonisin, trichothecenes and zearalenone in cereals. *Int J Mol Sci* 2008;9:2062-90.
  90. Pestka JJ, Zhou H, Moon Y, Chung YJ. Cellular and molecular mechanisms for immune modulation by deoxynivalenol and other trichothecenes: unraveling a paradox. *Toxicol Lett* 2004;153:61-73.
  91. International Agency for Research on Cancer (IARC). Toxins derived from *Fusarium graminearum*, *F. culmorum* and *F. crookwellense*. In: IARC Monographs on the evaluation of carcinogenic risks to humans, some naturally occurring substances: food items and constituents, heterocyclic aromatic amines and mycotoxins. Vol. 56. Lyon: IARC Press; 1993. p. 397-444.
  92. Schuhmacher-Wolz U, Heine K, Schneider K. Report on toxicity data on trichothecene mycotoxins HT-2 and T-2 toxins. CT/EFSA/CONTAM/2010/03 [displayed 31 October 2012]. Available at <http://www.efsa.europa.eu/en/scdocs/doc/65e.pdf>
  93. Riley RT, Wang E, Schroeder JJ, Smith ER, Plattner RD, Abbas H, Yoo HS, Merrill AH Jr. Evidence for disruption of sphingolipid metabolism as a contributing factor in the toxicity and carcinogenicity of fumonisins. *Nat Toxins* 1996;4:3-15.
  94. International Agency for Research on Cancer (IARC). Ochratoxin A. In: IARC Monographs on the evaluation of carcinogenic risks to humans, some naturally occurring substances: food items and constituents, heterocyclic aromatic amines and mycotoxins. Vol. 56. Lyon: IARC Press; 1993. p. 489-521.

**Sažetak**

## ŠTETNI UČINCI KOMBINIRANIH MIKOTOKSINA

U članku je prikazan pregled mikotoksinske kokontaminacije hrane u Hrvatskoj i susjednim zemljama te njihovih kombiniranih toksičnih učinaka na različitim eksperimentalnim modelima. Pritom su obuhvaćene studije interakcija koje uključuju fuzarijske mikotoksine, okratoksin A (OTA) i aflatoksin B<sub>1</sub> (AFB<sub>1</sub>). Nekoliko je takvih istraživanja napravljeno na temelju matematičko-statističkog modela, dok je većina studija primijenila jednostavnu statističku analizu koja omogućava usporedbu učinaka kombiniranih mikotoksina u odnosu na učinke pojedinačnih. Općenito, većina dosadašnjih studija pokazuje da kombinacije mikotoksina u biološkom sustavu imaju sinergistički ili barem aditivni učinak, što znači da su velik rizik za zdravlje ljudi i životinja.

**KLJUČNE RIJEČI:** *aflatoksin, citrinin, fumonizini, interakcije mikotoksina, okratoksin, trihoteceni, zearalenon*

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