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Research Article

Pathomorphological Changes in the Spleen of Turkey Broilers Challenged with Aflatoxin B1 Alone or Co-Administered with Mycotox NG -

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Summary

The aim of the present study was to evaluate the toxic effects of aflatoxin B₁ (AFB₁) on spleen morphology after its independent application or in combination with a mycosorbent (Mycotox NB) in turkey broilers. Experiments were carried out with 60 female turkey broilers 7-day age (meat TM strain) and divided into one control and five treatment groups (n = 10). Groups were as followed: Group I - control (fed standard feed according to the species and age of birds); Group II - feed by supplement with 0.5 g/ kg Mycotox NG, Group III - feed by supplementation with - 0.2 mg/ kg aflatoxin B₁, Group IV - feed supplementation with - 0.4 mg/ kg aflatoxin B₁, Group V - fed by supplementation with - with 0.2 mg/ kg aflatoxin B₁ and 0.5 g/ kg Mycotox NG and Group VI - fed by supplementation with - 0.4 mg/ kg aflatoxin B₁ and 0.5 g/ kg Mycotox NG. The duration of the experiments was 42 days. Specimens were collected from the spleen of birds for histological examination after euthanasia by cervical dislocation, fixed in 10% neutral formalin, dehydrated in alcohol series and embedded into paraffin. The 3 µm cross sections were cut on a microtome (Leica RM 2235) and stained with haematoxylin-eosin. Pathomorphological changes affected mainly the lymphatic tissue of germinative centres, whose number and size were reduced in birds from group IV and most cells exhibited degenerative (necrobiotic to necrotic) processes. In birds of group III, degenerative changes were less intensive (granular and vacuolar dystrophy) compared to the other groups. In birds of groups V and VI, the supplementation of toxin binder reduced partly the severity of morphological changes as congestion among lymphatic follicles in red and white pulp, slight granular and vacuolar dystrophy.

Keywords: Aflatoxin B₁; Turkey broilers; Spleen; Mycotox NG

INTRODUCTION

Aflatoxins are groups of closely related biologically active substances with minor differences in their chemical structure [1]. For the first time, aflatoxins were isolated more than 40 years ago in an outbreak of Turkey X disease Cullen and Newberne, [2] and rainbow trout cancer Blount, [3] after feeding diets comprising peanuts and cotton meal. These mycotoxins are secondary toxic metabolites produced by the genus *Aspergillus* (*Aspergillus flavus* and *Aspergillus parasiticus*) - coumarin derivatives containing a dihydrofurofuran moiety.

Aflatoxins are fluorescent compounds: under ultraviolet rays, aflatoxin B₁ (AFB₁) and aflatoxin B₂ (AFB₂) have a blue fluorescence while aflatoxins G₁ (AFG₁) and G₂ (AFG₂): yellow-green [4]. Aflatoxin B₁ (AFB₁) is the commonest metabolite in plant substrates and at the same time, the most toxic one. It is frequently encountered in cereal crops and peanut meal Gowda, et al. [5] and is the most potent liver carcinogen [6]. The toxicity of aflatoxin B₂ (AFB₂), G₁ (AFG₁) and G₂ (AFG₂) is 10%, 20% and 50% of that of AFB₁, respectively [7]. Aflatoxins are a group of almost 20 *Aspergillus* metabolites with similar chemical structure responsible for decay of plant substrates [8].

Aflatoxins are resistant to food processing and therefore could be found both in animal feeds and foodstuffs for direct consumption or processed foods, posing a risk for human health [9].

All bird species are sensitive to aflatoxin toxicity and although they do not receive relatively high concentrations with the feed to end lethally, low levels could be also deleterious after continuous intake. Growing birds, especially ducklings and turkey poultry are extremely sensitive to the toxic effects of aflatoxins. The total aflatoxin content in the feeds for poultry should not exceed 20 µg/ kg feed. Nevertheless, levels lower than 20 µg/ kg feed also induce lower resistance to diseases and stress [10]. The toxic effects of aflatoxins in domestic fowl are studied in detail with regard to their carcinogenic, teratogenic, mutagenic and growth inhibiting effects [11,12]. The occurring haematological, biochemical (decreased serum total protein, albumin, inorganic phosphorus, uric acid, total cholesterol and the values of haematocrit, red blood cell counts, mean corpuscular volume, haemoglobin, thrombocyte counts, percentage of monocyte counts; increased values of white blood cell and heterophil counts) Oguz, et al. [13], immunological (depression in anti-Brucella

abortus antibodies) Qureshi, et al. [14] and morphological (hydropic degeneration in liver; significantly reduced in size bursa of Fabricius and depletion of lymphoid cells from the follicles with necrosis and hyperplasia of mucosal epithelium, intraepithelial cysts oedema in the intraepithelial areas and heterophil infiltration; reduction in the size thymus and depletion of lymphocytes from the cortical and medullary areas; slight lymphoid depletion of periarteriolar lymphoid tussies in spleen; pale kidney with degeneration and/or necrosis of tubular epithelium [15].

The morphological changes in the spleen are characterized with lymphocytic degeneration, fatty dystrophy and haemorrhagic foci [16], congestive events in the red pulp, reduction of lymphocytes, vacuolar dystrophy [17], depletion of lymphoid cells, reticular cell hyperplasia, lymphocytolysis and increased germinal cell counts.

A number of strategies for detoxication of mycotoxin-contaminated feeds are proposed: physical separation, heat inactivation, irradiation, microbial degradation, treatment with chemicals. These methods of detoxication of aflatoxin-contaminated feeds are expensive [18]. The utilization of inert mycosorbents as hydrated calcium sodium aluminosilicate (HSCAS) [19], zeolites [20], bentonites [21], active charcoal [22] or bioproducts - live yeast cultures [23] are among the most extensively studied. Mycosorbents reduce the bioavailability of mycotoxins in blood and prevent their absorption in the intestines. Regardless of that, some of them reduce also the bioavailability of amino acids and/or minerals [24].

The aim of the present investigation was to evaluate the toxic effects of aflatoxin B₁ (AFB₁) on spleen morphology after its independent application or in combination with a mycosorbent (Mycotox NB).

MATERIAL AND METHODS

Experimental design

The experiment was performed with 60 7-day-old female turkey broilers (from the meat TM strain) randomly divided into six groups (n = 10).

All birds were fed standard feed according to the species and age, produced by a feed factory. The experimental design comprised: Group I - control; Group II - experimental, supplemented with 0.5 g/ kg Mycotox NG (micronised yeasts, montmorillonite, thymol); Group III - experimental, whose feed contained 0.2 mg/ kg aflatoxin

B₁; Group IV - experimental, whose feed contained 0.4 mg/ kg aflatoxin B₁; Group V - experimental, supplemented with 0.2 mg/ kg aflatoxin B₁ and 0.5 g/ kg Mycotox NG; Group VI - experimental, supplemented with 0.4 mg/ kg aflatoxin B₁ and 0.5 g/ kg Mycotox NG.

Aflatoxin B₁ used in this experiment was produced by *Aspergillus flavus*, 99% purity (Sigma-Aldrich, (Germany). All birds were housed under optimum microclimatic conditions as per [25].

After the trial, turkey poults from the control and experimental groups were euthanised via cervical dislocation according to [26]. Spleen specimens for histological study were fixed in 10% neutral formalin, dehydrated in ascending alcohol series and embedded in paraffin. Paraffin blocks were cut on a microtome Leica RM 2235 (Leica Biosystems Nussloch GmbH, Germany), with cross section thickness of 3 µm. They were stained with haematoxylin-eosin using routine histological techniques.

The experiment was approved by the Bulgarian Food Safety Agency - permit No 19218/06.11.2014. Statistical analysis was done with Statistica 6.0 (Windows) software, StatSoft, Inc. (USA, 1993) and ANOVA test. Data are presented as mean ± standard deviation (SD). The level of statistically significance was $p < 0.05$.

RESULTS

The relative weights of the spleen (g/100 g live weight) are presented in table 1. They were statistically significantly lower in birds from Groups III and IV (by 28.79% and 31.82%, respectively) compared to controls ($p < 0.001$). The addition of toxin binder to the feed of Groups V and VI prevented the reduction of relative spleen weights. There was no statistically significant difference between relative spleen weights of groups V and VI vs the control group ($p > 0.05$).

Pathomorphological studies

The most important pathomorphological changes in the spleen affected predominantly the lymph tissue of germinal centers. Turkey poults from the experimental group III supplemented with 0.2 mg/ kg AFB₁, exhibited mild degenerative changes comprising karyopyknosis, karyorrhexis and karyolysis of nuclei and rarefaction of lymphoid cells (Figure 1). Generalized hyperplasia of tunica intima of splenic arterioles could be detected (Figure 2).

In turkey poults from Group IV -, which received 0.4 mg/ kg AFB₁ in their feed, the degenerative changes in the spleen were rather more extensive. Lymph follicle rarefaction was enhanced, with necrotic foci in some areas (Figure 3). Occasionally, hemorrhagic foci could be seen in the splenic pulp. Red pulp hemosiderosis was observed as well (Figure 4). Along with these changes, sclerosis of the central artery was present (Figure 5).

The degenerative changes in Group V (0.2 mg/ kg AFB₁ and 0.5 g/kg Mycotox NG) were of lower intensity compared to those established in turkey poults from Groups III and IV. Morphological changes consisted mainly in congestive events in the red pulp and decreased density of lymphoid cells (Figure 6).

Turkeys from Group VI (0.4 mg/ kg AFB₁ + 0.5 g/ kg Mycotox NG) demonstrated mild lymphoid cell depletion with slight thickening of the intima of splenic arteries and occasionally, hemosiderosis (Figure 7).

There were no dystrophic changes in the spleen of control (Group I) and Mycotox NG supplemented (Group II) turkey poults.

DISCUSSION

Mycotoxinoses are diseases induced by feeding mould-contaminated feed ingredients. Aflatoxin is the prevailing and primary metabolite found in animal feeds [27]. Aflatoxins are a serious concern for poultry industry due to their high toxicity and common occurrence in compound poultry feeds [28,29]. In most cases, aflatoxin content in poultry feeds is higher than the maximally allowed amount of 20 µg. kg⁻¹ feed [23].

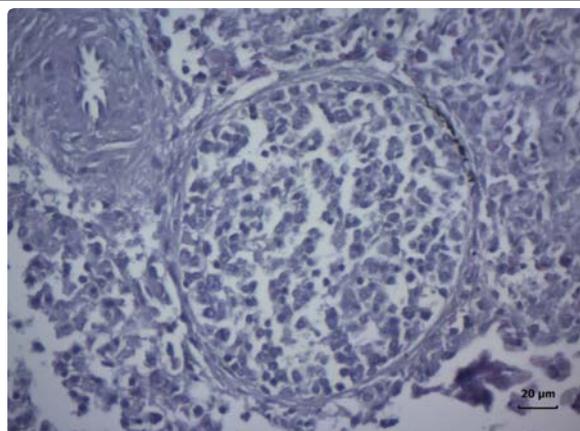


Figure 1: Weak to moderate degeneration - karyopyknosis and karyolysis of a splenic lymphatic follicle in a turkey poult treated with 0.2 mg/ kg AFB1 (Group III), H/E, Bar 20 µm

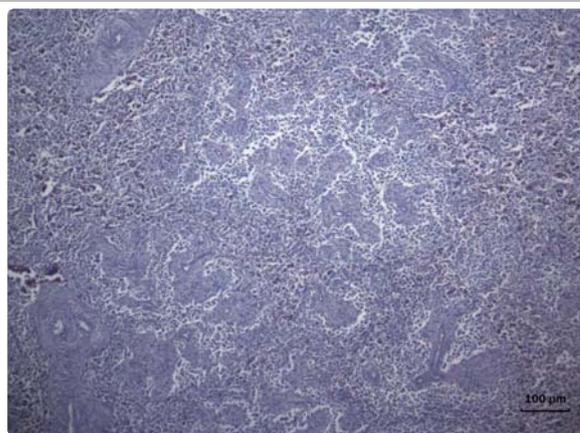


Figure 2: Hyperplasia of tunica intima of splenic arterioles in a turkey poult treated with 0.2 mg/ kg AFB1 (Group III), H/E, Bar 100 µm

Table 1: Effects of aflatoxin B₁ (AFB₁), alone or co-administered with Mycotox NG on the relative weight of the spleen in turkey poults.

Groups	I	II	III	IV	V	VI
	0.132 ± 0.021 ^a	0.136 ± 0.021 ^a	0.094 ± 0.093 ^b	0.09 ± 0.009 ^b	0.16 ± 0.008 ^a	0.15 ± 0.015 ^a

Different superscripts ^{a,b} indicates significant differences ($p < 0.05$ or more) between experimental and control groups ($P < 0.05$). The results are presented as mean ± SEM; n = 10 in each group.

Birds and fish are highly sensitive to the toxic impact of AFB₁, and react to low doses within the range 15-30 ppb [30]. The bioactivation of aflatoxins is done in hepatocytes through microsomal enzyme systems - cytochrome P450 (CYP450) where they are transformed into reactive aflatoxin-8,9-epoxide (AFBO), which is the primary most toxic metabolite. This reactive compound inhibits protein synthesis, causes liver damage, immunosuppressive effects and decreases productive performance. Domestic turkeys are highly sensitive to the toxic effects of AFB₁. In them, liver class alpha glutathione S-transferases (GSTA) are not capable to detoxify AFBO, which is probably the main reason for the high susceptibility of the species [30].

Sehu, et al. [31] reported that aflatoxins reduced the weight of immunocompetent organs (thymus, spleen, bursa of fabricius) in quails fed feed with total aflatoxin content of 2.5 mg/ kg. Compared to control group, the relative weight of the spleen in dietary treatments with AFB₁ only was lower. This reduction of immunocompetent organ weights was possibly due to necrosis and decreased density of lymphoid cells [32].

Aflatoxins induce immunosuppression in domestic fowl by altering the morphological structure of immunocompetent organs (thymus, spleen, bursa of Fabricius) [33]. Rats supplemented with

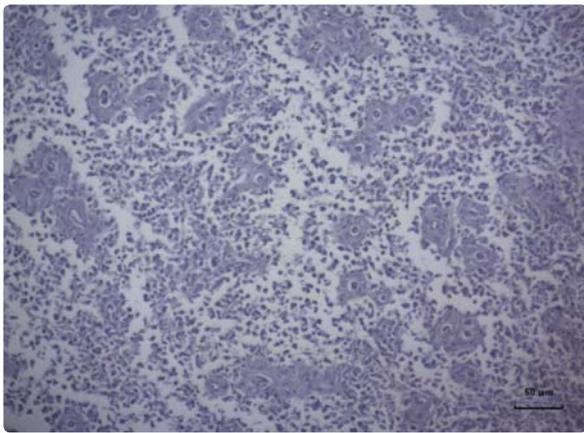


Figure 3: Necrotic foci in the spleen of a turkey poult treated with 0.4 mg/ kg AFB₁ (Group IV), H/E, Bar 50 µm.

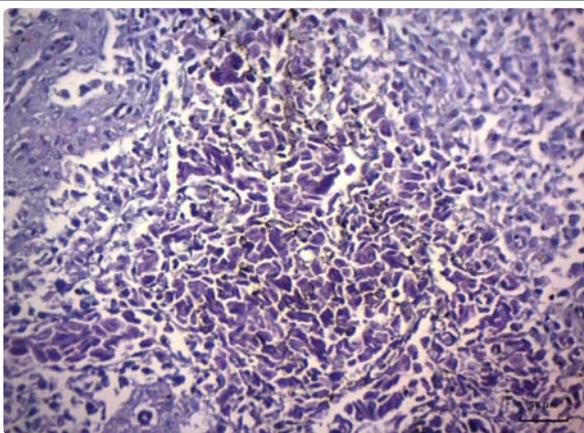


Figure 4: Haemosiderin granules in the spleen of a turkey poult treated with 0.4 mg/ kg AFB₁ (Group IV), H/E, Bar 50 µm.

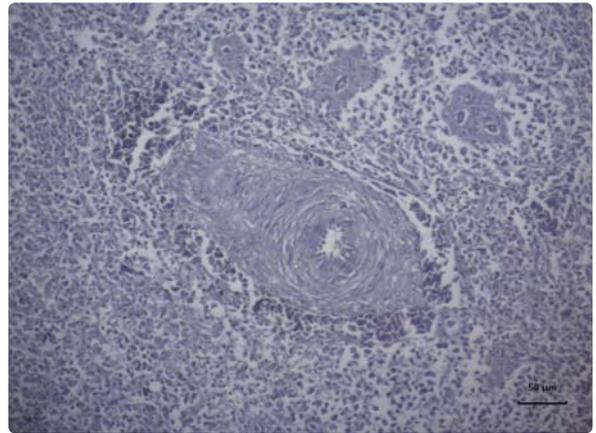


Figure 5: Central artery sclerosis in the spleen of a turkey poult treated with 0.4 mg/ kg AFB₁ (Group IV), H/E, Bar 50 µm.

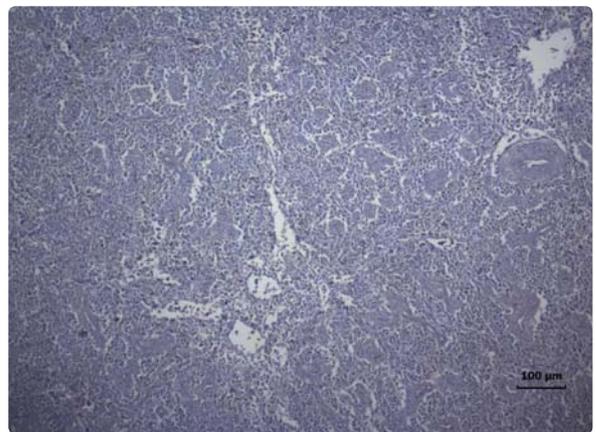


Figure 6: Red pulp congestion and lymphoid cell depletion in a turkey poult treated with 0.2 mg/ kg AFB₁ + Mycotox NG (Group V), H/E, Bar 100 µm.

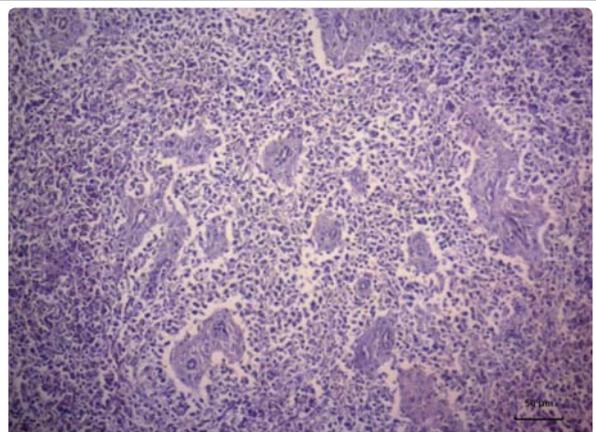


Figure 7: Mild lymphoid cell depletion with slight thickening of tunica intima of splenic arterioles in a turkey poult treated with 0.4 mg/kg AFB₁ + 0.5 g/ kg Mycotox NG (Group VI), H/E, Bar 50 µm.

1 mg/kg AFB₁ with the feed for 6 weeks exhibited degenerative and haemorrhagic foci in the splenic pulp, degenerated blood cells, megakaryocytes, irregular arrangement of leukocytes in the white pulp, haemosiderin granules, and congestion [34]. In



broiler chickens treated with 0.2 mg/ kg AFB₁, Sakhare, et al. [35] observed necrotic foci in germinative centers, central artery sclerosis, reduced number of lymphocytes, hyperplasia of tunica intima and destruction of elastic fibres. Decreased density of lymphoid cells in immunocompetent organs (bursa of Fabricius, thymus, spleen, caecal tonsils, Harderian glands) were found out in broiler chickens treated with citrinin and aflatoxin either alone or in combination [36]. Balachandran and Hashem, Mohamed, [37,38] demonstrated lower lymphoid cell density and necrotic changes in the bursa of Fabricius and the spleen, increased number of germinal centres and reticular hyperplasia in broiler chickens receiving 1 mg aflatoxin/ kg feed over 28 days. Observed splenic lesions also supported the immunotoxic and haemotoxic effects of aflatoxins [39,40]. Immunotoxic effects of aflatoxins are well studied in domestic poultry [33,41,42]. In previous studies of ours with broiler chickens Valchev, et al. [43] treated by supplementation of the feed with 0.5 mg/ kg AFB₁ or 0.8 mg/ kg AFB₁ dystrophic changes in lymphoid cells and reduced lymphoid cell density (lymph tissue reduction) were found out. Rats treated with 1 mg/ kg AFB₁ for 6 weeks exhibited lymphocyte degeneration, hemorrhages, degenerated blood cells, megakaryocytes, congestion in blood sinuses, necrotic foci, atrophy of the white pulp and irregular arrangement of leukocytes in the white pulp, [16]. The morphological structural changes demonstrated by this study were similar to changes in immunocompetent organs (thymus and bursa of Fabricius) reported by Kumar and Balachandran, [44] in broiler chickens treated with 1 mg/ kg AFB₁ over 28 days. It was demonstrated that the toxic effect of aflatoxins on lymphoid or hematopoietic organs resulted in rarefaction of lymphoid cell density in the white pulp and red pulp hemosiderosis [45].

Aflatoxins impair the synthesis of proteins by forming adducts with DNA, RNA and proteins [46], RNA synthesis inhibition, reduced activity of DNA-dependent RNA polymerase and degranulation of granulated endoplasmic reticulum [47] - mechanisms through which the structure of a number of tissues (liver, kidneys, skeletal muscles, heart, pancreas, immunocompetent organs) [48-50]. It is reported that aflatoxins enhance lipid peroxidation (LPO) and induce cellular damage resulting in alteration of normal morphological structure of parenchymal organs [51,52].

The addition of Mycotox NG to the diet of birds from Groups V and VI reduced the deleterious effects of AFB₁ on histopathological changes in the spleen of broiler chickens. Less severe histological lesions of immunocompetent organs were found out in broiler chickens after supplemented of aflatoxin-contaminated feed with natural zeolite (clinoptilolite) [53,41], herbo-mineral mycosorbent-Toxiroak [35] or of Mycotox NG [43]. The negative polarity of mycotoxins is attracted by the positive polarity of the toxin binder - a mechanism through which toxin are immobilized and eliminated from the organism of animals [54].

CONCLUSION

The tested AFB₁ amounts induced dose-dependent morphological alterations in the spleen (degenerative changes in lymph follicle, hyperplasia of tunica intima of splenic arterioles, rarefaction with necrotic foci), the relative weight of the spleen was lower compared to that of control birds. The addition of 0.5 g/ kg mycosorbent (Mycotox NG) to the ratio of Groups V and VI that contained either 0.2 mg/ kg or 0.4 mg/ kg AFB₁ partly reduced the deleterious effects of AFB₁ on spleen relative weight and the histological changes (congestive events decreased, density of lymphoid cells and thickening of the intima of splenic arteries and occasionally, hemosiderosis) in this organ.

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