Efficacy of Herbomineral Toxin Binder ‘Vilocym Z’ in Amelioration of Mixed Mycotoxicosis in Broilers

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Abstract: A study was conducted in 72 day old Vencobb broiler chicks to evaluate toxic effects of aflatoxin B1 and ochratoxin A and efficacy of herbomineral toxin binder product Vilocym Z (supplied by Ayurvet Ltd., Baddi, India) in preventing co-mycotoxicosis. Chicks were randomly divided into three groups (n = 24) and each group having three replicated of 8 chicks each. Group I served as healthy Control (C) and given standard basal ration and no treatment, Group T1 and T2 birds were fed standard basal diet and mycotoxicated with 100 ppb each of aflatoxin B1 and ochratoxin A from 0-21 days. Group T3 is not given any treatment and served as positive control; however, mycotoxicated group T1 was administered herbomineral toxin binder product Vilocym Z@1kg/tone of feed for 6 weeks. Mycotoxin adversely affected body weight gain, feed consumption, feed efficiency, haematobiochemical profile. However, supplementation of herbomineral toxin binder feed supplement has provided amelioration in mixed mycotoxicosis in broilers.

Key words: Aflatoxin, broiler, performance, ochratoxicosis, herbo-mineral, toxin-binder

INTRODUCTION

Aflatoxins are toxic secondary metabolites produced by fungi, namely Aspergillus spp. and Penicillium spp. High levels of aflatoxins have been recorded in ingredients of poultry feed soybean, sunflower, polished rice, cotton seed, etc. (Jand et al., 1995). The adverse effect of aflatoxins depends on age, species, nutritional status of the birds as well as dose and period in which it is consumed. Increased susceptibility of aflatoxicated chicks to infectious diseases indicates impaired immune responses. Aflatoxicosis leads to immunosuppression, characterized by decreased immune response (Baksch et al., 2000) and breakdown of the vascular immunity (Panisup et al., 1982). Similar effects of ochratoxin A with target organ kidney were summarized earlier by Marquardt and Frohlich (1992). Deleterious effect of aflatoxin could be overcome, or at least diminished, by adsorbents in rats (Abdel-Wahhab et al., 2002). Chemical adsorbents (Kubena et al., 1993), Levanoside hydrochloride (Kalorey, 1993), glucosemannan (Raju and Devegowda, 2000) as well as Growell (Godbole et al., 2001) have been attempted with varying degrees of success to reduce toxicity and impairment of immune response during aflatoxicosis in birds. In addition to this, another important mycotoxin is Ochratoxin (OTA), which are isolated from Aspergillus ochraceus but can also be produced by a species of Aspergillus and Penicillium species (Gibson et al., 1990). Of this group of isocumarins, only ochratoxin A has been naturally isolated from cereals and is the most toxic mycotoxin for birds. The natural occurrence of OTA in food and feedstuffs of plant and animal origin is common. OTA has been implicated in a diverse range of toxicological effects, including renal toxicity, mutagenicity, teratogenicity, neurotoxicity and immunotoxicity in both animals and man (O’Brien and Dietrich, 2005). OTA causes significant loss to poultry industry, intoxication of birds by ochratoxin results in reduced weight gain, impaired feed efficiency, reduced egg production and quality (Page et al., 1980). Use of adsorbents is of limited value in controlling ochratoxicosis in livestock (Marquardt and Frohlich, 1992; Santin et al., 2002). Stoev et al. (2000) and Kurkure et al. (2000) recently reported that 5% aqueous extract of artichoke and Curcuma Longa (Turmeric) powder at 0.5 g/kg feed reduces the toxic effect of ochratoxin A and aflatoxin B1 respectively, in chicks. Hence the present investigation was carried out to study the protective role and efficacy of herbal toxin binder product in broiler during induced combined aflatoxicosis and ochratoxicosis.

MATERIALS AND METHODS

Seventy two (72) day old broiler chicks were purchased and randomly divided into three identical groups (C, T1, T2) each comprising 24 chicks and each group was further comprising of 3 replicates of 8 chicks each. All the three groups were housed under identical managerial and environmental conditions in deep litter system from 0-6 weeks. Standard poultry feed free from aflatoxin and ochratoxin (basal ration) was offered

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all the three groups. The required quantity of ration for feeding to Control group C (negative control) was kept separately. The remaining feed was incorporated with 100 ppm of aflatoxin B1 and 100 ppm of ochratoxin A for feeding the birds belonging to groups T0 and T1 from 0-42 days. Chicks of group T0 was offered afla and ochratoxin contaminated feed without any mycotoxin binder product from 0-42 day. Treatment groups T1 was given mycotoxin binder product Vilocym Z @1 kg/tonne of feed from 0-42 days along with the afla and ochratoxicated feed from 0-42 days. Vilocym Z comprises of major herbs namely Phyllanthus emblica, Azadirachta indica, Citrullus colocynthis, Curcuma longa and many more herbs indicated for binding, inactivation and bio neutralization of multiple mycotoxins, maintenance and improvement in liver and kidney functions and for improved immune health and reduced oxidative stress in birds.

Production of aflatoxin B1: A known aflatoxin B1-producing strain of Aspergillus parasiticus (NRRL 3240) maintained on Sabouraud’s dextrose agar 2% (w/v) and aflatoxin B1 standard of 1 µg/mL, available at the Department of Microbiology, Nagpur Veterinary College, Nagpur, was used for production of aflatoxin and quantification of aflatoxin B1, respectively. The fungal spores were washed from the surface of agar slant with sterile Sabouraud’s Dextrose Broth (SDB) containing an equal amount of 0.1% Tween 80. The spore suspension was filtered through sterile muslin cloth and adjusted with SDB to a concentration of 1 x 10^6 spores/mL and was used as inoculum immediately. Two hundred and fifty g crushed soya DOC was sterilized in a 1 L conical flask and after cooling 25 mL of SDB was added to moisten the rice. One mL of the above mentioned inoculum was then added. It was then thoroughly mixed to ensure uniform distribution of spores and incubated at 28±1°C for 15 days. The flasks were shaken twice a day to break up clumps. After incubation, flasks were autoclaved at 10 Lbs for 5 m. The aflatoxin B1 was semiquantified according to Tapia (1985) using thin layer chromatography.

Production of ochratoxin A: Ochratoxin A (OA) was produced on crushed soya DOC as per the procedure described above, using a known ochratoxin A-producing strain of Aspergillus ochraceus (NRRL 3174) available at the Department of Microbiology, Nagpur Veterinary College, Nagpur. OA standard (3 µg/mL) was used for quantification of OA, according to Tapia (1985). Different parameters evaluated were growth, performance, haematological parameters, biochemical, enzymatic and gross pathology. Among growth promotion parameters, weekly record of body weight, feed consumption, Feed Conversion Ratio (FCR) was done and mean final body weight was also recorded. Blood samples were collected from five representative birds from each group twice during six weeks experimental period i.e. at the end of 3rd and 6th week to estimate haematobiochemical parameters. Haematological parameters included Haemoglobin (Hb), Packed Cell Volume (PCV), Total Erythrocytic Count (TEC), Total Leukocytic Count (TLC) and biochemical parameters included serum total proteins, albumin, globulin, lipid profile i.e. Total cholesterol, triglycerides, High Density Lipids (HDL), Low Density Lipids (LDL), Very Low Density Lipids (VLDL), serum creatinine, serum uric acid, SGOT, SGPT, tests were done using kits of Span Diagnostics Ltd., India. All the parameters were statistically analyzed by the method given by Snedecor and Cochran (1994).

RESULTS AND DISCUSSION

Growth and performance parameters: Average weekly body weight of broilers in various treatment groups is presented in Table 1. Gradual and significant (p<0.01) decrease in average body weight was observed in mycotoxin fed group T0. Significant (p<0.05) improvement and higher average body weight (1912 g) was observed in induced mycotoxicated groups treated with Vilocym Z (T1) in comparison to mycotoxicated and untreated group T0 (1753 g), during 1st, 3rd and 5th week of the experiment and found well comparable with the average body weight of birds of control group C (1952 g). Similar observations due to feeding of aflatoxin and ochratoxin were noticed earlier by Huff and Doerr (1981), Giambone et al. (1985), Raju and Devegowda (2000) and Stoew et al. (2000). There was significant (p<0.01) improvement in the body weight of treated group with herbal toxin binder product during induced mycotoxicosis. Earlier, Average weekly FCR of broiler is presented in Table 2. Significantly lower FCR was observed in prophylactically treated group T1 (1.92) during than untreated and mycotoxicated group T0 (2.15) at 6th week of experiment and found nearer to the FCR of birds of control group (1.91), indicating efficacy of the herbal toxin binder in ameliorating the toxic effects of the mycotoxin in the broilers.

Haematological parameters: Average haematological values of experimental broilers observed at 21st and 42nd day of experiment are presented in Table 3, respectively. Significant (p<0.01) reduction in values of Haemoglobin (Hb), PCV, TEC and TLC in mycotoxin fed group T0 was observed as compared to control group C during both periods of experiment. Significant (p<0.01) improvement in hematomical values were recorded in treatment group supplemented with Vilocym Z as compared to mycotoxin fed group T0 during the experiment and the values were well comparable with birds of control group C at both period of experiment. Significant reduction in Hb in broilers fed mycotoxin is in
correlation with the earlier findings of Doerr and Huff (1980) and Mani et al. (1993) on aflatoxicosis and Mohiuddin et al. (1993) and Ramadevi et al. (2000) on ochratoxicosis. Reduction in TEC and PCV due to feeding of aflatoxin (Singh et al., 1992) and ochratoxin (Doerr and Huff, 1980; Aved et al., 1991; Mohiuddin et al., 1993) was also reported earlier. Aved et al. (1991) and Mohiuddin et al. (1993) recorded a decrease in TLC due to induced aflatoxicosis and ochratoxicosis. The reduction in Hb concentration observed during mycotoxicosis could be due to reduced protein synthesis, as observed in the present study. Supplementation of polyherbal toxin binder showed improvement in various haematological parameters during induced mycotoxicosis.

**Biochemical parameters:** Average serum biochemical values of experimental broilers observed at 21st and 42nd day of age are presented in Table 4. Significant reduction in serum total protein, albumin and globulin was observed in mycotoxicated positive control group (Group T0) when compared to negative control group (Group C) on both 21st and 42nd day of experiment. The values of total protein, albumin and globulin were recorded to get normalized in prophylactically treated group with Vilocyzm Z and well comparable to healthy control and significantly higher than group T0 indicative of efficacy of polyherbal formulation in ameliorating the toxic effects of mycotoxicosis on liver and normalizing the serum values. Significantly higher values of serum total protein was found in the treated group T1 than group T0 and well comparable to the healthy chicks from control at both periods. Present findings are in agreement with Kaleory (1993), who recorded similar biochemical changes due to aflatoxin. Manning and Wyatt (1984), Ramadevi et al. (2000) and Stoev et al. (2000) reported decreased serum proteins during induced ochratoxicosis in broilers. Doerr and Huff (1980) and
Huff et al. (1992) reported reduction in serum total protein due to synergistic action of dietary aflatoxin and ochratoxin in chicks. In this experiment higher total serum protein, albumin and globulin values in polyherbal treated group T1, in contrast to untreated groups, showed restorative role of preparation as far as protein synthesis is concerned. Liver enzymes SGOT and SGPT were found to be significantly (p<0.01) elevated in induced combined mycotoxicosis (Group T0) when compared to negative control (Group C) at both the intervals. However, prophylactically treated group T1 supplemented with polyherbal toxin binder product showed significant (p<0.01) reduction in SGOT and SGPT levels than group T0 and found well comparable to birds of group C leading to normalization of liver during mycotoxicosis indicating efficacy of Vilocym Z in ameliorating the toxic effects of mycotoxin on liver and keeping the liver in healthy state. Elevation in values of liver marker enzymes (ALT and AST) has been reported at various levels of aflatoxins (Borisava et al., 1987; Raina et al., 1991) and ochratoxins (Gawale et al., 2009). Only serum total cholesterol level was significantly (p<0.05) decreased in mycotoxin fed group T1 while values of serum triglycerides, HDL, LDL and VLDL were found non significantly lowered when compared with control group C during 21st day of experiment. At 42nd
day of experiment, serum total cholesterol, triglycerides and VLDL levels were found significantly (p<0.01) lowered in mycotoxin fed group T0 in a comparison to control group, while serum values of HDL and LDL found non significantly lowered. Significant (p<0.01) improvement in serum cholesterol and non significantly higher levels of triglycerides, HDL, LDL and VLDL at 21st day of experiment in addition to significantly (p<0.01) increased levels of serum total cholesterol, triglycerides and VLDL along with non significant higher values of HDL and LDL at 42nd day of experiment were found in prophylactically treated groups with herbal toxin binder than mycotoxin fed group T0 and found well comparable group C indicating efficacy of herbal toxin binder in restoring the toxic effects of mycotoxin and normalizing the fat metabolism. Reduction in serum cholesterol during aflatoxicosis were reported earlier by Mani et al. (1993) and Vassan et al. (1998), likewise by Manning and Wyatt (1984), Ramadevi et al. (2000) and Stoev et al. (2000) during ochratoxicosis. The findings of present study are also in concomitance with those of Johri and Beura (2000) with Avsorb+ and Jindal et al. (1993) with
HSCAS (0.5%). Serum creatinine values were significantly (p<0.05 and p<0.01) higher in mycotoxin fed group T6 than control group during both the periods of observation. Supplementation of polyherbal toxin binder during mycotoxicosis significantly (p<0.01 and p<0.05) prevented a rise in values of serum creatinine in treated group than group T6 and found well comparable with serum creatinine values of control group C during both intervals of observation. Serum uric acid level in broilers was found to be significantly (p<0.01) elevated due to induced dietary mycotoxicosis in group T6 than control group. However, significant reduction in levels were recorded in treated group T1 receiving poly herbal toxin binder than group T6 during mycotoxicosis and found well comparable with the values of control group at both the interval. Present findings are in agreement with those of Manning and Wyatt (1984), Ramadevi et al. (2000), Doerr and Huff (1980) and Huff et al. (1992) in respect of ochratoxin and aflatoxin combination, respectively. The increase in serum creatinine and uric acid may be attributed to the nephrotoxic effect of ochratoxin, as evident in the present study, leading to renal dysfunctions. The findings in present study are also in corroborations with those reported by Sakhare et al. (2007) that feeding of Toxiroak® to mycotoxicated broilers significantly prevents a rise in the creatinine value, indicating its protective effect on kidney during mycotoxicosis.

Histopathology: Histopathology of liver of representative birds done on day 42nd revealed vacuolar degenerative changes and focal areas of necrosis in hepatocytes along with perportal necrosis in the mycotoxicated group T6 (Fig. 2) in contrast to normal orientation of hepatocytes in the liver section of healthy control group (C) birds (Fig. 1). Only mild congestion and haemorrhages were evident in the mycotoxicated and treated group T1 (Fig. 3). In the kidney of group T6 birds, cloudy swelling and severe glomeruli nephropathies along with multiple haemorrhages were observed in comparison to mild tubular degeneration and haemorrhages in group T1 (Fig. 4, 5). However, the architecture was normal of glomeruli and tubules in control group C.

Conclusion: There was a significant deleterious effect of mycotoxins on body mass, Feed Conversion Ratio (FCR), haematobiochemical parameters and body organs in broilers. Protection of changes in broilers supplemented with Vilocym Z @ 1 kg/tonne of feed for 0-42 days was recorded in terms of improving growth and performance parameters and normalizing the haematobiochemical profile in co-mycotoxicated broilers. The observed gross pathological changes on body organs, liver, kidney, spleen, bursa of fabricius and thymus of broilers were also alleviated by supplementation of Vilocym Z, as observed in the treatment group (T). It can be concluded that polyherbal toxin binder product ‘Vilocym Z’ is efficacious in ameliorating mixed mycotoxicosis in poultry.

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