INFLUENCE OF INCREASING FLUORIDE DOSE RATES ON SELECTED LIVER AND KIDNEY ENZYMES PROFILE IN DOMESTIC CHICKEN (Gallus domesticus)


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ABSTRACT

Fluoride has been considered to cause hepatic and renal tissue damages in animals and humans. The present study investigated the effect of varying concentrations of fluoride on hepatic and renal enzyme profile in domestic chicken (n=80). Chicken were distributed into 4 groups. Group A was kept unexposed while group B, C and D were exposed to 10, 20 and 30 μg/g body weight of NaF respectively on weekly basis for four weeks. Alkaline phosphatase (ALP), aspartate aminotransferase (AST), alanine amino-transferase (ALT) and bilirubin were determined as indicators of liver function test (LFT), while uric acid was as a parameter for renal function test (RFT). All LFT and RFT parameters showed high values (P< 0.05) after one, two three and four weeks in all groups. 579.4 ± 1.55, 355.0 ± 2.13, 246.2 ± 2.45 and 0.83 ± 1.46 were the ALP, AST, ALAT and bilirubin values for LFT and uric acid was 6.74 ± 2.92 in D group at the end of four weeks. All these results indicate the probability of severe effect on the physiology of the liver and kidneys in the exposed birds.

Key words: Fluoride intoxication, domestic chicken, LFT, RFT, bilirubin.

INTRODUCTION

Although fluoride intake is necessary for the development of teeth and body skeleton but requirements are in traces. However, a few recent studies indicated that more fluoride intake might cause toxic effects in animals and human being (Bhatnagar et al., 2011). Fluoride intake (fluorosis) usually occurs in two forms; endemic fluorosis, related to intake of drinking water with high fluoride contents (Li and Cao, 1994) and industrial fluorosis, related to exposure to air rich in fluoride contents (Czerwinski et al., 1988; Grandjean et al., 1985). Fluorosis has been found to cause severe side effects, not only to skeletal parts of the body (Finkelman et al., 1999; Wei, 1979) but also to the soft tissues like brain, liver, kidney and spinal cord (Guan et al., 1998; Wang et al., 2004).

Earlier work by Guan et al. (2000) reported that 50-80% of the absorbed fluorides are eliminated by the kidneys indicating the chances of kidney damages due to fluorosis. Liver is the main organ for fluoride detoxification and, therefore, is highly susceptible to the fluoride intoxication (Wang and Li, 2002). Various studies demonstrated that elevated levels of serum hepatic and renal enzymes have been found following fluoride intoxication indicating degenerative and inflammatory damages to the liver and kidney (Shivashankara et al., 2000; Wang and Li, 2002; Shashi and Thapar, 2001). Other histopathological findings include hyperplasia, dilatations of hepatic sinusoids (Kour et al., 1981), and accumulation of amorphous and crystalline bodies in the hepatocytes around the hepatic vein (Phillips et al., 1934), and in some rare cases the hepatocholan-gi carcinomain (Maurer et al., 1990). It has been found that fluoride intoxication induced hepatic hemangiosarcoma, hepatocellular adenoma and carcinoma, metastatic lung tumors, and Zymbal’s gland tumors in rats (Bogdanffy et al., 1995).

Commercial poultry production is being done extensively throughout the Pakistan. Further, management and production varies with the region which might be due to changes in the water and the presence of industrial wastes. Fluoride intoxication could be one of the intoxicants. Therefore, considering the growth of poultry sector and intensification of poultry farms around the urban areas, we hypothesized that higher levels of fluoride can affect the profile of liver and kidney enzymes. Selected serum liver enzymes were: alkaline phosphatase, aspartate aminotransferase, alanine aminotransferase, bilirubin and uric acid and for kidneys. Therefore, objective of current experiment was to evaluate the effect of fluoride intoxication on changes in liver and kidney enzymes of domestic chicken (Gallus domesticus).
MATERIALS AND METHODS

Eighty, a day old, domestic chickens (Gallus domesticus) were purchased from a local market. Chicks were reared till the age of four weeks before the start of experiment. The experimental protocols were approved by the ethical committee on animal research, GC University, Lahore (GCU).

Experimental Design: The birds were reared at the Animal House, Government College University, Lahore. After adaptability period, chicks were assigned one of the four dietary treatments namely A, B, C and D with 20 chicks per treatment. Group A served as a control group that was not treated with any fluoride intoxication; whereas, in group B, C and D, chicks were injected a weekly dose of NaF 10, 20, and 30 μg per gram of body weight, respectively. Chicks were sacrificed at the end of experiment and blood samples were collected.

Determination of Activities of Enzymes: Blood was collected from wing vein of chicken and then centrifuged at 5,000 rpm for five minutes to harvest the serum that were stored at - 20°C till further analysis. At the time of analyses, the serum samples were thawed and concentrations of alkaline phosphatase, aspartate aminotransferase, alanine aminotransferase, bilirubin and uric acid were determined using (MEDPHARM) commercial kits.

Data Analysis: Data, presented as mean ± S.D, were analyzed using one-way analysis of variance (ANOVA) and Tukey’s posthoc multiple comparison test. Data analysis was performed using statistical package SPSS (Version 13.0 SPSS Inc., Chicago, IL, USA) and results were declared significant at P < 0.05.

RESULTS AND DISCUSSION

When chicks were exposed to acute fluoride intoxication, higher (P<0.05) values of ALP were observed in all treatment groups. The ALP in control group was reported as 339.2 ± 1.44. The present study shows that ALP increased in an increased dose of fluoride manner, in group B ALP was 444.8 ± 2.32; whereas, 507.1 ± 1.54 and 579.4 ± 1.55 was observed ALP in group C and D. These results clearly indicate that the ALP was higher in those groups receiving the higher dose of fluoride intoxicant as shown in Table 1.

The serum AST also increased (P<0.05) with the increasing dose of fluoride. In control group, AST was 198.4 ± 2.44; whereas, in group B concentration was 268.2 ± 1.90. Highest values 310.9 ± 2.12 and 355.0 ± 2.13 of AST were observed in C and D groups respectively. Furthermore, serum ALT was 168.7 ± 1.55, 204.7 ± 4.18, 220.0 ± 3.56 and 246.2 ± 2.45 in the control B, C and D groups respectively shown (Table 1 & Fig1). The bilirubin level in control was 0.52 ± 0.04 U/L which increased in all treatment groups. The highest level of bilirubin (0.83 ± 1.46 U/L) was reported in the group D at the end of one month exposure to NaF as shown in Fig 2.

Increased serum bilirubin level indicates the impairment of liver functions as it is generally considered as a true test for proper liver functioning. It reflects the liver’s ability to take up and secrete bilirubin into the bile. The increase in ALP, AST, ALT and bilirubin was due to the degeneration and necrosis of liver cells. Shashi and Thapar (2001) reported that necrosis depends on the time of fluoride exposure. They showed that hepatic cell necrosis and other structural changes were more pronounced in the rabbits exposed to fluoride intoxication for three months than those exposed for one month. Similar study was conducted in human by Michael et al. (1996) to investigate the effect of high dose of fluoride intake from drinking water on some soft tissues and found that high levels of fluoride disturbed the normal ALT and AST values. The elevated ALP, AST, ALT and bilirubin is an indication of the impairment of liver functions. Most of the liver function enzymes were found abnormal in the children having skeletal fluorosis (Shivashankara et al., 2000). All these studies support our present findings of higher AST, ALP, ALT and bilirubin levels in the blood serum of chicken exposed to high doses of fluoride.

RFT parameter, uric acid also increased (P<0.05) in treatment groups. The uric acid level in the control was 3.63 ± 2.45; whereas, 4.54 ± 2.86, 5.55 ± 1.88 and 6.74 ± 2.92 (Table 1) were the reported values of uric acid after one month in group A, B, C and D respectively as shown in Fig 2. High uric acid indicates the impairment in the kidney functions as there are several evidences that hyperuricemia is the causative agent of kidney diseases (Sowers, 2004). High uric acid also aids in the development of other diseases like the albuminuria, microvascular disease, glomerulosclerosis, and tubulointerstitial fibrosis (Nakagawa et al., 2002). There are evidences that uric acid is a potent risk factor for kidneys in normal population as well as the population having kidney disease due to immunoglobulin A (IgA) nephropathy. A comparative study also indicate that hyperuricemia have more severe effects to the kidneys than proteinuria and make the kidneys less efficient and develop the kidney insufficiency (Sanchez-Lozada et al., 2005). High uric acid is a prediction of hypertension, metabolic syndrome (Masuo et al., 2003), diabetes (Nakanishi et al., 2003) and stroke (Bos et al., 2006). The present study provides us informations about the serum enzymes which are basic indicators of liver and kidney functioning at different fluoride exposure. It will be helpful to devise some strategies to protect the liver, kidneys and skeleton before they are fully damaged and get irreversible loss.
Table 1. Effect of increasing levels of fluoride on liver and kidney serum enzymes of domestic chicken.

<table>
<thead>
<tr>
<th>Treatment Groups</th>
<th>A</th>
<th>B</th>
<th>C</th>
<th>D</th>
</tr>
</thead>
<tbody>
<tr>
<td>ALP</td>
<td>339.2±1.14</td>
<td>444.8±2.32</td>
<td>507.1±1.54</td>
<td>579.4±1.55</td>
</tr>
<tr>
<td>AST</td>
<td>198.4±2.44</td>
<td>268.2±1.90</td>
<td>310.9±2.12</td>
<td>355.0±2.13</td>
</tr>
<tr>
<td>ALT</td>
<td>168.7±1.55</td>
<td>204.7±4.18</td>
<td>220.0±3.56</td>
<td>246.2±2.45</td>
</tr>
<tr>
<td>Bilirubin</td>
<td>0.52±3.20</td>
<td>0.61±3.21</td>
<td>0.65±2.58</td>
<td>0.83±1.46</td>
</tr>
<tr>
<td>Uric Acid</td>
<td>3.63±2.45</td>
<td>4.54±2.86</td>
<td>5.55±1.88</td>
<td>6.74±2.92</td>
</tr>
</tbody>
</table>

abcd within a row means for each variable without a common superscript differs significantly at P < 0.05.

1 Treatments: Group A (control) without fluoride, whereas, Group B, C and D, chicks were injected a weekly dose of NaF 10, 20, and 30 µg per gram of body weight, respectively.

REFERENCES


subsequent weight gain and blood pressure elevation. Hypertension 42(4):474-480.


