Original Research Article

Influence of Dietary Supplementary Betaine Hydrochloride on Blood Biochemical Profile in Broilers

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Abstract

The study was initiated to examine the effect of dietary supplementation of betaine hydrochloride (betaine HCl) on Serum biochemistry in broiler chicken. The experimental feed was formulated according to BIS (1992) specifications and to the control ration (T1), feed grade betaine HCl at 250, 500, 750 ppm was added respectively to form different rations T2, T3 and T4 for different treatment groups. The birds in each group were maintained on their respective ration throughout the experimental period of six weeks. Four birds from each treatment was randomly selected at six weeks of age to study the haematological parameters. All serum parameters were within the normal range for species. Supplementation of betaine HCl in different levels did not improve the serum calcium and inorganic phosphorus. The serum total protein and albumin level as well as serum uric acid were similar (P>0.05) in all treatment groups. There was significant decrease in serum lipid levels triglycerides (P< 0.01) and low density lipoproteins (P< 0.05) among treatment groups but serum total cholesterol, high density lipoproteins, aspartate aminotransferase (AST) and alanine aminotransferase (ALT) were similar (P>0.05) in all treatment groups.

Keywords
Broiler birds, Betaine hydrochloride, Hematocrit, Biochemical profile

Introduction

Betaine is a naturally occurring amino acid derivative found in microorganisms, plants and animals. Many foods such as wheat, shellfish, spinach and sugar beets contain significant amount of betaine. Metabolic role of betaine in the body is primarily as the methyl group donor and it is also an osmolyte that assists in cellular water homeostasis. Betaine is a zwitterionic quaternary ammonium compound that is also known as trimethylglycine, glycine betaine, lycine, and oxyneurine. It is a methyl derivative of the amino acid glycine with a formula of \((\text{CH}_3)_3\text{N}^+\text{CH}_2\text{COO}^-\) and a molecular weight of 117.2, and it has been characterized as a methylamine because of its 3 chemically reactive methyl groups (Yancey et al., 1982 and Kidd et al., 1997). Zeisel et al. (1994)
noted that choline act as a methyl donor for the formation of methionine from homocysteine, by being oxidized to betaine.

**Materials and Methods**

One hundred and ninety two, day-old Vencobb – 400 strain commercial broiler chicks were used as the experimental birds. The birds were allotted to four dietary treatment groups, with four replications of 12 chicks each randomly in a completely randomized design. The experimental feed (in mash form) was formulated using corn and soybean meal as per BIS (1992) specifications. To the control ration (T1), feed grade betaine HCl was added at 250, 500 and 750 ppm to formulate rations T2, T3 and T4 respectively, taking special care for proper mixing of betaine HCl. No growth promoting substance were added to any rations.

Approximately 10 ml of blood was collected during slaughter at 42nd day from four birds each belonging to treatment groups fed rations T1, T2, T3 and T4 from jugular vein in sterilized glass tubes for serological analysis. The collected blood was allowed to clot and centrifuged for 10 minutes at 2000 rpm to separate the serum and stored in deep freezer at -20°C for further analysis. Serum samples were analyzed for total cholesterol, triglycerides, HDL cholesterol, total protein, albumin and globulin by using Semi Automated Biochemical Analyser (Master T). The LDL cholesterol was calculated by Friedewald equation.

Mineral profile serum calcium and phosphorus was estimated by reports of Moorehead and Biggs, 1974 and Miller et al. (1994). Total protein and albumin concentration quantified by standard procedure (Bromocresol green method) (Lowry et al., 1951 and Johnson et al., 1999) respectively. Estimation of lipid profiles total cholesterol (Richmond, 1973), serum triglycerides (Bucolo and David, 1973), HDL (Seigler and Wu, 1981) and LDL (Friedewaldet et al., 1972) done as method given in parenthesis of researcher. Excretory metabolic profile uric acid concentration by Uricase-PAP method (Lypo CHEK kit) and liver enzymatic profile Serum aspartate aminotransferase (SGOT or AST) and alanine aminotransferase (SGPT or ALT) contents were measured by using Liqui CHEK kit (M/s. Agappe Diagnostics Limited, Ernakulam, Kerala).

The data collected on various parameters were statistically analyzed as per the methods of Snedecor and Cochran (1994) and the means of different experimental groups were also tested by using Duncan’s Multiple Range Test (DMRT) in SPSS Version 20.0.

**Results and Discussion**

The mean values obtained for serum minerals in experimental birds fed four dietary treatments T1, T2, T3 and T4 were 10.98, 10.54, 10.61 and 10.70 mg/dl for calcium and 6.27, 6.33, 6.37, 6.29 mg/ dl for inorganic phosphorus are presented in Table 1.

The serum biochemical parameters of experimental birds maintained on four dietary treatments T1, T2, T3 and T4 were 5.15, 4.74, 4.89 and 4.98 g/ dl for total proteins and 2.87, 2.47, 2.57 and 2.75 g/ dl for albumin and 2.27, 2.27, 2.32 and 2.23 g/ dl for globulin, 1.27, 1.12, 1.12 and 1.27 g/ dl for A/G ratio, 6.00, 6.75, 6.73 and 6.73 g/ dl for uric acid respectively are depicted in Table 2.

The serum lipid profile (mg/ dl) of experimental birds belonging to the groups T1, T2, T3 and T4 were 23.11, 18.68, 14.95 and 13.96 for triglycerides, 72.71, 69.16, 66.57 and 64.46 for total cholesterol, 29.66, 27.72, 31.31 and 33.63 for HDL cholesterol and
38.44, 37.70, 32.27 and 28.03 for LDL cholesterol respectively are represented in the Table 3. The liver enzymes (U/L) of experimental birds belonging to treatment groups T₁, T₂, T₃ and T₄ were 255.06, 241.96, 254.85 and 246.49 for aspartate amino transferase (SGOT) and 13.72, 13.04, 12.62 and 13.03 for alanine amino transferase (SGPT) respectively are presented in Table 4.

**Table 1** Serum calcium and inorganic phosphorus level of birds maintained on four experimental rations, (mg / dl)

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Treatments¹</th>
<th></th>
<th></th>
<th></th>
<th>F value</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>T₁</td>
<td>T₂</td>
<td>T₃</td>
<td>T₄</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Calcium</td>
<td>10.98 ±0.57</td>
<td>10.54 ±0.76</td>
<td>10.61 ±0.50</td>
<td>10.70 ±0.36</td>
<td>0.118ns</td>
<td>0.948</td>
</tr>
<tr>
<td>Inorganic Phosphorus</td>
<td>6.27 ±0.16</td>
<td>6.33 ±0.10</td>
<td>6.37 ±0.19</td>
<td>6.29 ±0.19</td>
<td>0.066ns</td>
<td>0.977</td>
</tr>
</tbody>
</table>

¹Mean of 4 observations with SE
ns-non significant (P>0.05)

**Table 2** Serum total protein, albumin, globulin, uric acid concentration (mg / dl) and albumin/globulin ratio (%) of birds maintained on four experimental rations

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Treatments¹</th>
<th></th>
<th></th>
<th></th>
<th>F value</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total protein</td>
<td>5.15 ±0.19</td>
<td>4.74 ±0.15</td>
<td>4.89 ±0.18</td>
<td>4.98 ±0.11</td>
<td>1.142ns</td>
<td>0.372</td>
</tr>
<tr>
<td>Albumin</td>
<td>2.87 ±0.08</td>
<td>2.47 ±0.22</td>
<td>2.57 ±0.19</td>
<td>2.75 ±0.12</td>
<td>1.259ns</td>
<td>0.332</td>
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<tr>
<td>Globulin</td>
<td>2.27 ±0.12</td>
<td>2.27 ±0.16</td>
<td>2.32 ±0.10</td>
<td>2.23 ±0.17</td>
<td>0.600ns</td>
<td>0.980</td>
</tr>
<tr>
<td>Albumin/globulin ratio</td>
<td>1.27 ±0.05</td>
<td>1.12 ±0.15</td>
<td>1.12 ±0.11</td>
<td>1.27 ±0.15</td>
<td>0.495ns</td>
<td>0.692</td>
</tr>
<tr>
<td>Uric acid</td>
<td>6.00 ±0.35</td>
<td>6.75 ±0.18</td>
<td>6.73 ±0.16</td>
<td>6.73 ±0.11</td>
<td>2.813ns</td>
<td>0.084</td>
</tr>
</tbody>
</table>

¹Mean of 4 observations with SE
Table 3: Serum lipid profile of birds maintained on four experimental rations, (mg / dl)

<table>
<thead>
<tr>
<th>Lipid profile</th>
<th>Treatments</th>
<th>T1</th>
<th>T2</th>
<th>T3</th>
<th>T4</th>
<th>F value</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Triglycerides</td>
<td></td>
<td>23.11±1.28</td>
<td>18.68±0.26</td>
<td>14.95±1.14</td>
<td>13.96±1.14</td>
<td>16.055**</td>
<td>0.000</td>
</tr>
<tr>
<td>Total cholesterol</td>
<td></td>
<td>72.71±2.42</td>
<td>69.16±2.86</td>
<td>66.57±3.79</td>
<td>64.46±1.16</td>
<td>1.702ns</td>
<td>0.220</td>
</tr>
<tr>
<td>High density lipoprotein</td>
<td></td>
<td>29.66±2.02</td>
<td>27.72±0.72</td>
<td>31.31±1.00</td>
<td>33.63±1.43</td>
<td>3.297ns</td>
<td>0.058</td>
</tr>
<tr>
<td>Low density lipoprotein</td>
<td></td>
<td>38.44±1.95</td>
<td>37.70±3.32</td>
<td>32.27±2.89</td>
<td>28.03±0.87</td>
<td>4.007*</td>
<td>0.034</td>
</tr>
</tbody>
</table>

1 Mean of 4 observations with SE
ns – non significant (P>0.05), ** significant at 0.01 level (P< 0.01)
* Means bearing different superscripts within same column differ significantly (P< 0.05)

Table 4: Serum aspartate amino transferase (SGOT) and alanine amino transferase (SGPT) levels of birds maintained on four experimental rations, (U/L)

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Treatments</th>
<th>T1</th>
<th>T2</th>
<th>T3</th>
<th>T4</th>
<th>F value</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Aspartate amino transferase (SGOT)</td>
<td></td>
<td>255.06±11.57</td>
<td>241.96±5.49</td>
<td>254.85±24.18</td>
<td>246.49±11.97</td>
<td>0.187ns</td>
<td>0.903</td>
</tr>
<tr>
<td>Alanine amino transferase (SGPT)</td>
<td></td>
<td>13.72±0.68</td>
<td>13.04±0.57</td>
<td>12.62±0.16</td>
<td>13.03±0.23</td>
<td>0.963ns</td>
<td>0.442</td>
</tr>
</tbody>
</table>

1 Mean of 4 observations with SE
ns – non significant (P>0.05)

The values of serum calcium and phosphorus concentration of broilers in the present study were within the normal range reported for the species and comparable with the values reported by Silva et al. (2007) and no difference (P<0.05) in serum calcium and phosphorus concentration among various treatment groups fed different levels of betaine HCl supplementation, and the values were within the normal range prescribed for the species.

Similarly Attia et al. (2005) and Konca et al. (2008) could not observe any significant effect on the serum protein levels with betaine supplementation in the broiler diet.

On the other hand Hassan et al. (2005), El-Husseiny et al. (2007) and Rao et al. (2011) in slow growing chicken and in the broiler chicken, respectively reported that betaine supplementation caused significantly higher serum protein levels while reduction in serum...
uric acid levels was reported by Zhan et al. (2006) in birds fed methionine deficient broiler diet supplemented with betaine supplementation.

The result on serum triglyceride revealed significant (P≤0.01) difference between groups fed four treatment rations. T1 (control) rations fed birds had significantly higher serum triglyceride than those fed rations T2, T3 and T4, but those fed T3 and T4 rations recorded lowest triglycerides and significantly differed from those fed T1 and T2 rations.

Statistical analysis of data on serum total cholesterol and HDL cholesterol revealed no difference between treatment groups. The treatment group fed T4 rations had significantly lower (P≤0.05) serum LDL cholesterol than the all other treatment groups but at the same time there was no difference between treatment groups fed T1, T2 and T3 rations. This is in agreement with the results of Jahanian and Rahmani (2008) who reported that betaine supplementation as a replacement for choline in broiler diets did not affect plasma levels of cholesterol and low density lipoproteins (LDL) but, resulted in significant decrease in plasma triglycerides and very low density lipoproteins and significant increase in high density lipoproteins.

El-Husseiny et al., (2007) indicated elevated levels of total lipids with increasing levels of betaine at 0.5, 0.75 or 1.0 g / kg of broiler diet but was contradictory with report of Maghoul et al. (2009) and Baghaei et al. (2011) who reported no significant effect on blood parameters like cholestrol, triglycerides, LDL, VLDL and HDL with betaine supplementation by replacement of choline and low methionine diet, respectively.

On statistical analysis of data on the above liver enzymes revealed no significant difference (p>0.05) between treatment groups. Attia et al. (2005) and Konca et al. (2008) in their study found that serum alanine amino transferase level in blood of birds was decreased when they were fed with diets supplemented with betaine.

There was no significant difference between treatment groups in serum minerals, protein and uric acid excretory profile. As per mentioned results suggest that there is no role of betaine in the above standard profile. (Table 1 & 2)

The serum lipid profile (mg/ dl) of experimental birds fed rations T1, T2, T3 and T4 were 23.11, 18.68, 14.95 and 13.96 for triglycerides, 72.71, 69.16, 66.57 and 64.46 for total cholesterol, 29.66, 27.72, 31.31 and 33.63 for HDL cholesterol and 38.44, 37.70, 32.27 and 28.03 for LDL cholesterol, respectively.

On statistical analysis, it was found that the triglycerides level of birds supplemented with betaine HCl at 750 ppm were significantly lower (P<0.01) than the unsupplemented group. The LDL cholesterol level in serum was significantly lower (P<0.05) for birds fed with betaine HCl at 750 ppm than the unsupplemented group. However Statistical analysis of data on serum total cholesterol and HDL cholesterol revealed no significant difference (P> 0.05) between treatment groups. There was no significant difference (P> 0.05) among treatment groups for SGOT and SGPT.

It is concluded clearly the supplementation of betaine HCl at 750 ppm in the diet decreased the serum triglycerides, serum LDL cholesterol and breast muscle cholesterol content, hence, the supplementation of betaine HCl can be exploited the betaine involvement in lipid metabolism. There is a real challenge of betaine HCl to decrease the both blood and breast muscle cholesterol.
References


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