Potency of garlic juice supplementation on some physiological and immunological aspects of broilers exposed to heat stress

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Abstract

The current study aimed to investigate the effect of garlic juice supplementation with broiler drinking water on performance, immune response, lipid profile, blood picture and activity of liver enzymes under heat stress. For this purpose 100 four days aged broiler checks (Ross) were allocated into control (C) and three treatment groups (25 each). All group checks were housed at 5 ºC above normal ambient temperature during the experiment. Treatment groups (T1, T2 and T3) checks were supplemented with 0.5, 1 and 1.5 ml/L, respectively, with garlic juice in drinking water (6 hrs a day). Food and water were supplemented ad libitum. Body weights were monitored every five days and blood samples were obtained at 25 days for assessment of blood picture and antibody titers after vaccination, total cholesterol and triglycerides concentrations and activity levels of alanine aminotransferase (ALT), aspartate aminotransferase (AST) and alkaline phosphatase (ALP) were estimated. At 30 days age, T3 group broiler checks reported higher body weights than other groups. Total number of white and red blood cells, hemoglobin concentration, hematocrit levels and ratio of heterocytes to lymphocytes increased significantly in treated groups, whereas significant decline of cholesterol and triglycerides concentrations as well as ALT and AST levels were shown in T3 group broiler checks. Significant increase was shown in the levels of antibody titers against both Newcastle (ND) and infectious bursal disease (IBD) vaccines. In conclusion, treatment of broiler checks early with garlic juice increase broiler performance and immune response against vaccination program and may be protect the broilers from heat stress.

Keywords: Garlic juice, Broilers, Immune response, Heat stress, Body weight, ALT, AST

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Introduction

In poultry farm, there is increasing insistence to use antibiotics as food additive as growth promoter, increase performance of broilers and for treatment of many bacterial diseases (1). However, the public health is affected due to the low doses of antibiotics which have been used as food additive in animal production farm, where this addition could develop bacterial resistance to the antibiotics in humans (2). Therefore, since 2006 European Union restricted the use of antibiotics as additive in animal (3). Thus, decreased performance and increased incidence of diseases in poultry population were produced as a result to antibiotics restriction in animal food. Many researchers have been used growth promoters of natural origin like organic acids, enzymes, essential oils of some plants, probiotics and yeast cultures to provide a new, safe and cheap growth promoter to recompense the use of antibiotics in animal food (4,5).

Garlic (Allium sativum), a member of family Alliaceae which was used as a spice, has antibacterial, anti-inflammatory, antioxidant and antifungal effects (6,7). The major medical impact effects of garlic appear to be due to the sulfur containing compounds, minerals and enzymes. The sulfur compounds in garlic are of two types, the S-alkylcysteine sulfoxides and the γ-glutamyl-s-alkylcysteines, which are presented in equal quantities. The most important and presented in high quantity in garlic sulfur compound is alliin (8). Garlic preparation by crushing or mincing resulting in activation of alliin by allinase which is amino acid present in intact garlic to produce allicin which is the main thiosulfate compound in the garlic and has a half life 16 hours at 25°C or 2.5 days when kept as juice or crushed, also, pH less than 3.5 or high environmental temperature make the allinase enzyme inactive and one minute in microwave is sufficient to inactivate this enzyme which is responsible for thiosulfate conversion (9).

In broilers, it was mention that garlic, can be used as feed additive, increase chicken growth and feed conversion ratio and improve vitality Tollba and Hassan (10). Amagase et al. (11) and Demir et al. (12) reported that modulation of poultry performance and carcass merits can be improved by adding garlic powder to the poultry diets. Lewis et al. (13) showed that addition of plant extracts to broilers’ diet has some effects on performance and microbial activity of intestinal tract but, none of them were significant. Moreover, Cho et al. (14) concluded that allicin has immune-stimulatory effect on birds. Demir et al. (12) attributed the strong stimulating effect of garlic to the immune system of broilers primarily to the bioactive components of garlic together with sulphur containing compounds such as alliin and allicin.

Sanjeev et al. (15) suggested that garlic powder should be added to the poultry diets because they find this addition improve growth and increase poultry tolerance to the stress of environmental sources. Garlic extract (allicin) supplementation at 50 mg/kg has an economic importance by increase feed conversion ratio, improve blood constituents parameters and chicken performance if it is used at recommended dose (16).

The present study was conducted to investigate the impact of fresh garlic juice supplementation with drinking water on some physiological and immunological aspects of broiler chicks exposed to heat stress.

Materials and methods

This study was carried out at the animal house of the College of Agriculture, University of Sumer, Iraq during the period extended from October 10, 2016 to February 1, 2017.

Preparation of the extract

Garlic was purchased from local market of Al-Refaa city. Garlic juice was prepared daily by rinsing suitable amount of the peeled garlic bulb in a ream with equal amount of distilled water quantity w/v and the resulted juice was incubate in 37 °C overnight then filtrated and the aqueous part was used directly with drinking water of treated groups (17).

Broiler checks

One hundred unsexed one day old Ross chicks were used in this experiment. Broiler checks were housed in ventilated clean cages (120*100*40 cm length, widen and height respectively) 25 bird per cage, and the temperature was 35 °C using electrical heater with automatic thermostat. Food and water were supplied ad-libitum. The chicks were vaccinated against most common infectious diseases in Iraq, Newcastle, avian Influenza and Infectious Bursal disease. Commercial starter and grower diets (compose
from corn, soybean, multivitamin, minerals, Di-calcium, Methionine, Lysine, choline chloride, and vegetable oil with crude protein 23% and digestible energy 3010 Kcal/kg for starter and crude protein 21% and 3100 Kcal/kg for grower) from Al-Waha Co., Iraq was used as basal diet. Food and water were supplied ad-libitum.

Experimental design
One hundred unsexed one day old Ross chicks were vaccinated against most common infectious diseases Newcastle, avian Influenza and Infectious Bursal disease (Intervet, Holland). The experimental periods were extended for 30 days. The broiler checks were individually weighed and randomly allocated into four groups (25 chicks each). All group checks were housed at 5 °C above normal ambient temperature during the experiment. First group broiler checks were fed on basal diet and served as control group. Second (T1), third (T2) and fourth (T3) groups were fed on the basal diet and And received drinking water contains garlic juice daily for 6 hours at dose 0.5, 1 and 1.5 ml/l respectively. Broiler checks were weighed initially and every 5 days interval. Wing blood samples were obtained at the end of experiment for assessment of total RBC, Hb and differential WBC count (18). ALT, AST, ALP (19), total cholesterol (20) and Triglyceride (21) concentrations were assessed using spectrophotometric methods. Immune response was measured by ELISA technique according to method described by Trinder (22) and Doumas et al. (23), respectively.

Statistical analysis
The results were expressed as mean ± SD. One way analysis of variance (ANOVA1) and Newman-Keuls were used to compare between experimental groups. Difference at a level of p<0.05 was considered as significant. Statistical analysis was performed using the GraphPad Prism version 5 (GraphPad Software, Inc. California, USA).

Results
The result in table (1) showed significant increase in body weight of T3 group checks compared with other groups which showed insignificant (p≥0.05) differences when compared between each other. The insignificant elevation started earlier at day 5 but it became significant (p<0.05) at day 20 of broiler age.

The results showed that the total number of WBC and RBC counts of T3 group checks increased significantly (p<0.05) compared with other groups (table 2). Also T2 group checks recorded higher number than T1 and control group checks, which was significantly lower than that of T1. Hemoglobin (Hb) concentrations of T3 and T2 group checks showed insignificant (p≥0.05) difference between each other, but they were significantly (p<0.05) higher than control and T1 group checks. In T3 group checks, the hematocrit (PCV) results showed significant (p<0.05) increase than other groups, while T2 group checks recorded the lowest value among experimental groups, whereas T1 and control group checks recorded insignificant (p≥0.05) difference between each other. In all treated group checks, the ratio between heterophils and lymphocytes showed significant (p<0.05) decrease compared with that of control, however in T3 group, the lowest value was recorded.

Table (3) showed significant (p<0.05) decrease in cholesterol levels of treated groups compared to that of control, also triglyceride concentration showed significant (p<0.05) decrease in T3 group among experimental groups. Total protein levels showed insignificantly (p≥0.05) difference among experimental groups, however, albumen concentration was significantly (p<0.05) increase in treated groups compared to that of control group.

The results in the present study showed significant (p<0.05) decrease in the activities of ALT, AST and ALP in all treated groups compared to that of control group (table 4). On the other hand, the results of immune response measured by ELISA showed significant (p<0.05) increase in the antibody titers against ND and IBD in T3 and T2 group checks compared to that of T1 and control group checks, which showed insignificant (p≥0.05) difference between each other (table 5).

Table 1: Body weight changes (gm) in control and garlic juice treated broiler checks

<table>
<thead>
<tr>
<th>Age (days)</th>
<th>C</th>
<th>T1</th>
<th>T2</th>
<th>T3</th>
</tr>
</thead>
<tbody>
<tr>
<td>5</td>
<td>134±5 a</td>
<td>133±4 a</td>
<td>135±3 a</td>
<td>136±5 a</td>
</tr>
<tr>
<td>10</td>
<td>245±4 a</td>
<td>246±4 a</td>
<td>250±6 a</td>
<td>256±8 a</td>
</tr>
<tr>
<td>15</td>
<td>530±6 a</td>
<td>532±8 a</td>
<td>537±10 a</td>
<td>540±15 a</td>
</tr>
<tr>
<td>20</td>
<td>854±10 c</td>
<td>870±10b</td>
<td>875±13 b</td>
<td>910±18 a</td>
</tr>
<tr>
<td>25</td>
<td>1126±20 c</td>
<td>1187±11b</td>
<td>1185±18b</td>
<td>1270±25a</td>
</tr>
<tr>
<td>30</td>
<td>1750±25b</td>
<td>1760±20b</td>
<td>1791±30b</td>
<td>1910±20a</td>
</tr>
</tbody>
</table>

Values were expressed as mean ± SD (n=15 of broiler chicks). Different small letters refer to significant differences (p<0.05) between groups for each period, C (control broiler checks): supplemented with water and food without any addition, T1 (treatment group-1): supplemented with 0.5 ml of garlic juice/L of drinking water daily for 6 hr a day, T2 (treatment group-2): supplemented with 1 ml of garlic juice/L of drinking water daily for 6 hr a day, T3 (treatment group-3): supplemented with 1.5 ml garlic juice/L of drinking water daily for 6 hr a day.
Table 2: Blood parameters changes in control and garlic juice treated broiler checks

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Groups</th>
<th>C</th>
<th>T1</th>
<th>T2</th>
<th>T3</th>
</tr>
</thead>
<tbody>
<tr>
<td>WBC (1000/ml)</td>
<td>24±0.3</td>
<td>24.3±0.2</td>
<td>24.8±0.3</td>
<td>25±0.1</td>
<td></td>
</tr>
<tr>
<td>RBC (10*12)L</td>
<td>2.36±0.2</td>
<td>2.37±0.2</td>
<td>2.57±0.2</td>
<td>2.59±0.2</td>
<td></td>
</tr>
<tr>
<td>Hb (g/dl)</td>
<td>10.5±1.1</td>
<td>11.4±1.1</td>
<td>9.9±1.1</td>
<td>11.4±1.1</td>
<td></td>
</tr>
<tr>
<td>g/dl</td>
<td>0.01 b</td>
<td>0.02 a</td>
<td>0.01 c</td>
<td>0.02 a</td>
<td></td>
</tr>
<tr>
<td>HCT</td>
<td>27.3±1.1</td>
<td>27.7±1.1</td>
<td>26±1.1</td>
<td>29.9±1.1</td>
<td></td>
</tr>
<tr>
<td>H/L ratio</td>
<td>0.27±1.1</td>
<td>0.23±1.1</td>
<td>0.23±1.1</td>
<td>0.21±1.1</td>
<td></td>
</tr>
</tbody>
</table>

Values were expressed as mean ± SD (n=15 of broiler chicks), Different small letters refer to significant differences (p<0.05) between groups for each period, C (control broiler checks): supplemented with water and food without any addition, T1 (treatment group-1): supplemented with 0.5 ml of garlic juice/L of drinking water daily for 6 hr a day, T2 (treatment group-2): supplemented with 1 ml of garlic juice/L of drinking water daily for 6 hr a day, T3 (treatment group-3): supplemented with 1.5 ml garlic juice/L of drinking water daily for 6 hr a day.

Table 3: Lipid profile changes in control and garlic juice treated broiler checks

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Groups</th>
<th>C</th>
<th>T1</th>
<th>T2</th>
<th>T3</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cholesterol mg/100ml</td>
<td>200±6 a</td>
<td>135±5 b</td>
<td>130±3 b</td>
<td>124±2 c</td>
<td></td>
</tr>
<tr>
<td>Triglycerides mg/100ml</td>
<td>183±3 a</td>
<td>182±2 a</td>
<td>182±3 a</td>
<td>167±4 b</td>
<td></td>
</tr>
<tr>
<td>Total protein g/dl</td>
<td>5.6±1 a</td>
<td>5.5±1 a</td>
<td>5.7±1 a</td>
<td>5.7±1 a</td>
<td></td>
</tr>
<tr>
<td>Albumin g/dl</td>
<td>3.8±1 b</td>
<td>4.9±1 a</td>
<td>4.8±1 a</td>
<td>4.9±1 a</td>
<td></td>
</tr>
</tbody>
</table>

Values were expressed as mean ± SD (n=15 of broiler chicks), Different small letters refer to significant differences (p<0.05) between groups for each period, C (control broiler checks): supplemented with water and food without any addition, T1 (treatment group-1): supplemented with 0.5 ml of garlic juice/L of drinking water daily for 6 hr a day, T2 (treatment group-2): supplemented with 1 ml of garlic juice/L of drinking water daily for 6 hr a day, T3 (treatment group-3): supplemented with 1.5 ml garlic juice/L of drinking water daily for 6 hr a day.

Table 4: Liver enzyme activities in control and garlic juice treated broiler checks

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Groups</th>
<th>C</th>
<th>T1</th>
<th>T2</th>
<th>T3</th>
</tr>
</thead>
<tbody>
<tr>
<td>AST µ/L</td>
<td>9±0.5 a</td>
<td>6±0.5 b</td>
<td>7±0.5 b</td>
<td>5±0.5 c</td>
<td></td>
</tr>
<tr>
<td>ALT µ/L</td>
<td>29±2 a</td>
<td>30±3 a</td>
<td>32±2 a</td>
<td>33±4 a</td>
<td></td>
</tr>
<tr>
<td>ALK µ/L</td>
<td>1767±11 a</td>
<td>899±8 b</td>
<td>681±10 c</td>
<td>461±9 d</td>
<td></td>
</tr>
</tbody>
</table>

Values were expressed as mean ± SD (n=15 of broiler chicks), Different small letters refer to significant differences (p<0.05) between groups for each period, C (control broiler checks): supplemented with water and food without any addition, T1 (treatment group-1): supplemented with 0.5 ml of garlic juice/L of drinking water daily for 6 hr a day, T2 (treatment group-2): supplemented with 1 ml of garlic juice/L of drinking water daily for 6 hr a day, T3 (treatment group-3): supplemented with 1.5 ml garlic juice/L of drinking water daily for 6 hr a day.

Table 5: Immune response changes in control and garlic juice treated broiler checks

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Groups</th>
<th>C</th>
<th>T1</th>
<th>T2</th>
<th>T3</th>
</tr>
</thead>
<tbody>
<tr>
<td>ND.</td>
<td>868±15 c</td>
<td>850±20 b</td>
<td>1316±20 b</td>
<td>1460±13 a</td>
<td></td>
</tr>
<tr>
<td>IBD</td>
<td>990±12 c</td>
<td>1030±15 b</td>
<td>1190±25 a</td>
<td>1135±35 a</td>
<td></td>
</tr>
</tbody>
</table>

Values were expressed as mean ± SD (n=15 of broiler chicks), Different small letters refer to significant differences (p<0.05) between groups for each period, C (control broiler checks): supplemented with water and food without any addition, T1 (treatment group-1): supplemented with 0.5 ml of garlic juice/L of drinking water daily for 6 hr a day, T2 (treatment group-2): supplemented with 1 ml of garlic juice/L of drinking water daily for 6 hr a day, T3 (treatment group-3): supplemented with 1.5 ml garlic juice/L of drinking water daily for 6 hr a day.

Discussion

The results of present study were in agreement with other studies (16,17,24-26) who found that supplementation of garlic to the broiler food has a positive effect on body weight. Also, this finding disagreed with (27) who found that the adding of garlic powder to the broiler food at different levels has no significant (p≥0.05) effect on the body weight of the chicks. The positive role of allicin on the performance of intestinal flora may be the cause of improving digestion and feed conversion ratio (28) in addition to the antibacterial effect of garlic on different types of pathogenic bacteria which is indirectly increase...
feed conversion ratio (29). The role of garlic acid in the enhancement of enzymes activity of pancreas which will provide a suitable environment for food absorption may be another cause for this result (30).

The present findings were in harmony with the results obtained by Oleforuh-Okelah et al. (31) who found that treatment with garlic and ginger in birds will significantly increase WBC, RBC, PCV and Hb, also, Elnagar et al. (32) found that the adding of garlic extract to the birds diet produce significant increase in the number of RBC. Therefore, it can be suggested that the increase in body weight may be due to the improving in oxygen carrying capacity of this constituents of the blood resulting in maximal availability of nutrients to the chicks. Jamroz et al. (33) suggested that the increase in the number of WBC is due to the bacteria static effect of garlic extract especially on *Salmonella* spp and *E-coli* which lead to produce large number of WBC due to abundance of low pathogenic antigen.

Moreover, the improvement in the chick immune system may be due to the immunostimulatory effect of garlic juice which cause significant decrease in the H/L ratio especially that shown in T3 group checks. This result was in agreement with that of Eidi and Iraqui (34) and Mohamed et al. (16) who observed significant increase in the number of lymphocyte in the broiler chicks supplied with garlic extract which could be the cause of decreased H/L ratio.

The results of lipid profile, shown in the present study, were in agreement with that of previous studies (16,33-36) who found that garlic supplementation has a hypocholesterolemic effect in treated birds. Garlic exhibit this effect may be due to its possible role in the inhibition of most important enzymes in cholesterol and lipid synthesis such as cholesterol 7α-hydroxylase, fatty acid synthetase and hepatic 3-hydroxy-3-methylglutaryl co-enzyme A reductase (15).

Many researchers reported that garlic or its extract supplementation has a protective effect on hepatocytes which will cause significant (p≤0.05) in the activity of liver enzymes (15,16,37). Generally the lowest enzyme activity was recorded in serum of broiler chicks of group T3. These results can be attributed to garlic juice supplementation which act as stabilizer to liver cell membrane and prevent the harmful effect of the deleterious agent and free radicals.

During vaccination, when the antigen enters the body, the macrophages ingest the antigen and then display a portion of the antigen on its membrane, which stimulates the B-lymphocytes to produce the antibodies. The macrophages also produce interleukin-1, Interferon γ and α, which stimulate B lymphocytes to produce antibodies. T-lymphocytes are differentiated and divided to affect T cells and helper T cells, which meet the antigen presented on the macrophage membrane. As a result of IL-1 stimulation, Interleuken-2 (IL-2) stimulates the response of B-lymphocytes, which produce antibody type IgM, which converts them into B-lymphocytes producing IgG antibodies (38). Garlic has an important role in this mechanism by stimulating division and differentiation of lymphocyte and macrophage, also, activate phagocytosis, IL-2 and γ interferon production (39,40). These results were in agreement with that of Mohamed et al. (16) who found that dietary supplementation of garlic improved phagocytosis and immune response.

In conclusion garlic juice supplementation to broiler drinking water at a dose of 1.5 ml/L daily for 6 hours a day for 30 days will increase body weight, immune response, protect liver cells against harmful effect of Heat stress and improve blood picture constituent.

### References


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