Use of feed additives in animal agriculture: gut response in broilers

1Ofongo - Abule, R. T. S. and 2Etebu, E.
1Department of Animal Science, Niger Delta University
2Molecular Genetics/Microbiology Research Unit, Department of Biological Sciences, Niger Delta University, Wilberforce Island, Bayelsa State – Nigeria

?Corresponding author: tariruth@live.de; phone: 08158683316; 08038827764

Abstract

Feed additives such as enzymes, probiotics, prebiotics and acidifiers are few examples of suitable alternatives to in –feed antibiotics in animal agriculture due to the ban of antibiotic use. However, the Nigerian poultry industry still experiences cases of antibiotic use. The gut benefits of alternatives to antibiotics often culminate in improved growth performance but there are or may be certain gut responses that attribute to the observed performance of poultry birds. In order to ascertain this, an experiment was conducted to evaluate the effect of antibiotic administration and enzyme supplementation on gut pH and bacteria counts. One hundred and fifty day-old broiler chicks were randomly distributed to three dietary treatments having five replicates and 10 birds per replicate. The experiment lasted forty – two (42) days and was arranged as a completely randomized design. A maize-soybean meal diet not supplemented with antibiotic or enzyme served as the control. Birds fed diet II had antibiotic (Dicoxin plus ®) administered to them. Birds fed diet III had their diets supplemented with enzyme (Roxazyme G2G®). Results indicated pH of the crop to be significantly (P<0.01) reduced by feed additive. The pH of the Ileum was least (5.65) under antibiotic administration but was not significantly (p>0.05) different that of birds fed control and enzyme supplemented diet. Gizzard pH was acidic but not significantly (P>0.05) across all the treatments. Lactobacillus and Coliform counts were significant (P<0.05) affected by feed additive addition. It can be concluded that feed additive impact on gut pH can influence the type and population of bacteria present in the gastrointestinal tract.

Keywords: gut pH, antibiotics, enzymes, gut bacteria, broilers

Introduction

The micro flora and digestive physiological parameters such as intestinal enzymes, pH, viscosity, histo - morphological structures of intestine plays a major role in production performance of broilers. Broiler chickens are reared up to 35 – 42 days of age in intensive system, where the birds are subjected to many types of stress that can result in reduced performance. To alleviate these types of stress, antimicrobials have been used as feed supplement for more than 50 years in poultry feed. Most of the antibiotic growth promoters act by modifying the intestinal flora, which are associated with poor health and reduced performance of animals (Bedford, 2000). Digesta pH is one of the major gastrointestinal environment influences on nutrient biodiversity (Shafey et al., 1991). Accurate determination of digesta pH in broilers could act as a tool to indicate potential for optimum gut health and hence maximize nutrient absorption (Morgan et al., 2012). Lower digesta pH in the gut is associated with a reduction in growth and colonization by pathogenic organisms thus permitting greater partitioning of nutrient for optimal growth and nutrient utilization. Consequently, lower gut pH could be associated with higher weight gain Dono et al. (2014). Lower pH in the intestine
inhibits the growth of especially pathogenic organism and favors growth of beneficial organism as reported by Andil et al. (2011). Removing digesta appears to negatively affect the accuracy of digesta pH reading (Morgan et al., 2012). Previous investigations using mainly culture-dependent approach showed that majority of Lactobacilli, Enterococci and Clostridia were found in the gut of broiler birds (Salanitro et al., 1978; Barnes, 1979; Mead, 1989; Engberge et al., 2000). The microflora is believed to protect against intestinal colonization by pathogens and are primarily responsible for degrading the copious amount of mucus produced by goblet cells in the intestine. Many factors can affect the composition of the avian bacterial community such as diet, age, antibiotic administration and infection of pathogenic organisms (Lu et al., 2003). Bacteria within the gut microbial community interact with each other as well as their host (De-Ageli et al., 2006; Kelly et al., 2005). Microorganisms can directly interact with the lining of the gut and immunological status of birds (Torok et al., 2007). The pH concentration of digesta collections from different segment of the gut of broilers was observed that most organic acid used in feed and drinking water are absorbed by the upper gastrointestinal segment i.e. crop, pro – ventriculus and gizzard and only a little portion of organic acid get to the lower digestive tract – caecum (Hummel et al., 1993). In the past years, a great deal of interest has been generated on the evaluation of the alternative means for manipulation of gastrointestinal micro flora in livestock production. Hinton et al., (1990) explained that the crop contains some microbes which produce lactic acid that helps to decrease crop pH. This study was designed to determine the effect of antibiotics and enzyme supplementation on gut pH and also to determine the intestinal microbial counts in broilers as affected by gut pH.

Materials and methods

Animal experiment
A total number of 150 one-day old (ANAC 2000) broiler chicks were utilized in this study. The chicks were brooded together for seven days at a temperature of 32°C. The birds were randomly distributed into three dietary treatments having five replicates of 10 birds per replicate. The diet consisted of maize – soybean meal (M-SBM) as main ingredient which served as control diet. Birds fed diet 2 was given the same diet with the control but were administered antibiotics at an inclusion rate of 100gm/160 litres in drinking water. The third diet was also the same as the control diet but was supplemented with a non – starch polysaccharide degrading enzyme (Roxazyme G2G ®). The birds were fed ad libitum and the experiment lasted 35 days.

Gut pH determination
On day 35, two birds per replicate were slaughtered by cervical dislocation to enable collection of digesta content of the crop, gizzard, ileum and both ceca. Digesta collected was utilized for pH determination with a pH meter (Model: HANNA Instrument – Hi 9024-microcomputer pH meter). The experimental diet was collected for each treatment and set aside for proximate analysis according to AOAC (1990).

Determination of gut micro flora
Sodium Chloride (NaCl), de Man Rogosa and Sharp (MRS) agar and McConkey agar was prepared and sterilized for 121°C in an autoclave, and timed for 15 minutes. The NaCl and agars were removed after sterilization and allowed to cool. The NaCl (9ml) was poured into six test tubes
respectively and 1gm wet weight of digesta collected from the ileum was added into the first test tube for serial dilution, a sterilized pipette was used to mix properly. The diluents in the fourth, fifth and sixth test tubes were plated out on MRS and McConkey agar in triplicates respectively. The petri dishes were kept upside down in an incubator (30°C on MRS agar for 48 hours to determine *Lactobacillus* and 37°C on McConkey agar for 24 hours to determine *Coliform*). Colony growth was counted and values obtained were log transformed before carrying out statistical analysis using SPSS volume 17 and significant means separated using duncans multiple range test.

<table>
<thead>
<tr>
<th>Feed ingredient</th>
<th>M - SBM</th>
<th>M – SBM + AB</th>
<th>M – SBM + EZ</th>
</tr>
</thead>
<tbody>
<tr>
<td>Maize</td>
<td>55</td>
<td>55</td>
<td>55</td>
</tr>
<tr>
<td>SBM</td>
<td>32.5</td>
<td>32.5</td>
<td>32.5</td>
</tr>
<tr>
<td>Fish meal</td>
<td>4.0</td>
<td>4.0</td>
<td>4.0</td>
</tr>
<tr>
<td>Cassava starch</td>
<td>4.2</td>
<td>4.2</td>
<td>4.2</td>
</tr>
<tr>
<td>Constant ingredients*</td>
<td>4.3</td>
<td>4.3</td>
<td>4.3</td>
</tr>
<tr>
<td>Total</td>
<td>100</td>
<td>100</td>
<td>100</td>
</tr>
</tbody>
</table>

Analysed nutrient composition (g/kgDM)

<table>
<thead>
<tr>
<th></th>
<th>M - SBM</th>
<th>M – SBM + AB</th>
<th>M – SBM + EZ</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dry matter (grams)</td>
<td>739.5</td>
<td>733.5</td>
<td>651.5</td>
</tr>
<tr>
<td>Ash</td>
<td>254.2</td>
<td>203.1</td>
<td>170.4</td>
</tr>
<tr>
<td>Protein</td>
<td>238.0</td>
<td>229.0</td>
<td>256.3</td>
</tr>
<tr>
<td>Ether extract</td>
<td>58.1</td>
<td>62.7</td>
<td>49.1</td>
</tr>
<tr>
<td>Crude fibre</td>
<td>64.9</td>
<td>73.6</td>
<td>70.6</td>
</tr>
</tbody>
</table>

*: Bone meal: 2.1kg; Oyster shell: 1.0kg; Vitamin/premix: 0.25kg, DL-Methionine: 0.15kg, Common salt 0.3kg, TiO₂: 0.5kg, SBM: soybean meal

**Results and discussion**

**Nutrient composition**

Nutrient composition of the experimental diets is presented in Table 1. Although all three experimental diets were similar but there was disparity in analyzed nutrient composition especially crude protein and ash concentration. The ash concentration was least in enzyme supplemented diet (170.4g/kgDM) but high in the control diet (254.2g/kgDM). However, the reverse was the case with regards to crude protein concentration which was least in the diet which antibiotic was administered (229.0g/kgDM) but high in the enzyme supplemented diet (256.3g/kgDM). Dry matter concentration was least in M – SBM +EZ diet (651.5g) compared to the control (739.4g) and antibiotic (M – SBM +AB) administration (733.5g).

**Gut pH**

Gut pH as affected by antibiotic administration and enzyme supplementation is presented in Table 2. Antibiotic administration and enzyme supplementation had significant (p<0.01) on pH of the crop. Values recorded were more acidic with a value of 4.83 recorded for antibiotic administration and 5.09 for enzyme supplementation.

<table>
<thead>
<tr>
<th>Gut Section</th>
<th>M – SBM</th>
<th>M – SBM +AB</th>
<th>M – SBM +EZ</th>
<th>SEM</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Crop</td>
<td>6.02b</td>
<td>4.83a</td>
<td>5.09a</td>
<td>0.04</td>
<td>0.008**</td>
</tr>
<tr>
<td>Gizzard</td>
<td>5.59</td>
<td>5.45</td>
<td>4.91</td>
<td>0.20</td>
<td>0.375ns</td>
</tr>
<tr>
<td>Ileum</td>
<td>6.46</td>
<td>5.65</td>
<td>6.10</td>
<td>0.18</td>
<td>0.215ns</td>
</tr>
<tr>
<td>Caecum</td>
<td>6.66</td>
<td>6.59</td>
<td>6.52</td>
<td>0.07</td>
<td>0.770ns</td>
</tr>
</tbody>
</table>

ab: means along the same row with different superscripts are significantly different (p<0.05).

**: 1% level of significant difference.
Gut pH gradually increased from highly acidic to near neutral from the crop to the caecum in birds administered antibiotics. However, this was not the case in the gut of broilers fed the control and enzyme supplemented diets. The pH value dropped in the gizzard in broilers fed the control diet from 6.02 in the crop to 5.59 and 5.09 in the crop to 4.91 in broilers fed enzyme supplemented diet. In all, pH values recorded for the respective gut segments in broilers fed enzyme supplemented and antibiotics administered birds was lower compared to the control but was not significantly different (p>0.05) across the treatments with regards to values recorded in the gizzard, ileum and caecum.

**Gut micro flora counts**

Table 3 showed that enzyme supplementation significantly (p<0.05) increased *Lactobacillus* counts in the ileum while antibiotic administration significantly (p<0.05) reduced *Lactobacillus* counts. It was observed that *Coliform* counts was significantly (p<0.01) higher in the ileum of birds fed enzyme supplemented diet (8.27) compared to the control (7.81) and antibiotic administration (7.67) which recorded significantly (p<0.01) lower values respectively.

Enzymes have tendency to increase crude protein of a diet because certain protein attached to carbohydrate molecules are released when such carbohydrate molecules are hydrolyzed by enzymes (Ofongo – Abule and Ohimain, 2016). According to Dono et al. (2014), lower digesta pH was associated with birds of high body weight irrespective of the phase (starter or grower or finisher). The high weight group in that study recorded a significantly lower proventriculus (gizzard) pH compared to a low weight group. This was also the case in this study. Birds fed enzyme supplemented diet had the least proventriculus pH and a significantly high weight gain (unpublished data).

A lower crop pH was observed due to antibiotic administration and enzyme supplementation. This is in line with an earlier report by Hinton et al. (1990) which stated that the crop contains some microbes (mostly *Lactobacillus*) which produce lactic acid that ultimately helps to decrease pH of the crop. However, the report of Hummel et al. (1993) differed from this in that the author reported an increase in crop pH. The methodology of pH determination may play a role in gut pH values recorded as stated by Morgan et al. (2012). Ofongo – Abule et al. (2016); who reported an acidic pH in the gut of broilers as resulting from the effect of diet type and enzyme supplementation. The authors recorded a gradual increase in gut pH from crop to ileum and eventually caecum. This trend was also observed in the current study in exception of pH value recorded in the gizzard of birds fed enzyme supplemented diet which was numerically low (4.91) or more acidic compared to values recorded in control and antibiotic administration. Furthermore, the report of Olukosi and Dono (2014) indicated that phytobiotic

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**Table 3: Effect of antibiotics administration and enzyme supplementation on gut microflora counts isolated from the ileum of broilers**

<table>
<thead>
<tr>
<th>Bacteria</th>
<th>M-SBM</th>
<th>M-SBM+AB</th>
<th>M-SBM+EZ</th>
<th>SEM</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Lactobacillus</em> spp</td>
<td>7.58&lt;sup&gt;b&lt;/sup&gt;</td>
<td>7.33&lt;sup&gt;b&lt;/sup&gt;</td>
<td>7.76&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.06</td>
<td>0.034&lt;sup&gt;*&lt;/sup&gt;</td>
</tr>
<tr>
<td><em>Coliform</em></td>
<td>7.81&lt;sup&gt;b&lt;/sup&gt;</td>
<td>7.67&lt;sup&gt;b&lt;/sup&gt;</td>
<td>8.27&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.06</td>
<td>0.003&lt;sup&gt;**&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

<sup>ab</sup>: means along the same row with different superscripts are significantly different (p<0.05).
<sup>*</sup>: 5% level of significant, <sup>**</sup>: 1% level of significant difference.
supplementation (turmeric and garlic meal) at 10g/kg of diet significantly reduced digesta pH in the crop, proventriculus and caeca but had no effect at the jejunum. Invariably digesta pH response may differ with additive supplemented to diets of similar ingredient composition. Although in – feed antibiotic use is banned in certain countries, however gut micro flora response to antibiotic use may differ when compared to other available alternatives. Gut micro flora response in this study indicated a significant \((p < 0.05)\) increase of \textit{Lactobacillus spp} in ileum of birds fed enzyme supplemented diet. However, that recorded for the control diet was higher than the value recorded in birds administered antibiotic but was not significantly different \((p>0.05)\). Based on pH values recorded in the ileum, it would have been expected that \textit{lactobacillus} counts would be higher in antibiotic administered diet than control or enzyme supplemented diet. According to Gadd (1997), antibiotics affect micro flora by altering the metabolism of microorganisms and suppressing microbial growth in the gut. This was also stated by Jones and Ricket (2003) that antibiotics decrease microbial load in the GIT and improve weight gain and feed conversion ratio thereby making more nutrients available to the host. According to Ohimain and Ofongo (2013) the stimulation of \textit{Lactobacillus} by a diet containing wheat offal with or without enzyme supplementation coincided with the reduction in \textit{coliform} and \textit{Escherichia coli} population. This observation was not the case in the current study but the least count of \textit{coliform} was recorded in antibiotic treatment. The individual variability in body weight and nutrient utilization in grower and finisher phases of broilers can be partly explained by variability in jejuna and caecal pH (Dono \textit{et al.}, 2014). Also this may be related to the type of microbes colonizing the gastro intestinal tract which will ultimately affect bird performance.

**Conclusion**

The study showed that antibiotic administration and enzyme supplementation lowers gut pH while \textit{Lactobacillus} count was increased by enzyme supplementation.

**References**


Shafey, T. M., Mcdonald, M. W. And


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