

Studies on Effect of Mannan oligosaccharides, β -glucan and their Combination on Immune Status and Litter Condition of Broilers Reared under Hot Climatic Condition

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Abstract

An experiment was conducted to evaluate the immune status litter condition of broilers by supplementation of mannan oligosaccharides (MOS), β -glucan and their combination. 320 one day old straight run Vencobb400 chicks were weighed and distributed randomly into four treatment groups viz, A, B, C and D with four replicates consists of 20 chicks in each replicate. Treatment group A was serving as control without supplementation of MOS and β -glucans. Supplementation of β -glucans @5g/100kg and MOS@ 50g/100kg of feed was done in treatment group B and C respectively. The treatment D was a combination of treatment B and C. The results revealed non-significant influence of supplementation of MOS, β -glucans and their combination on Ranikhet Disease (RD) titre at 21st and 42nd days of age. There were non-significant differences with pH and the litter moisture. It may be concluded that the supplementation of β -glucans, MOS and their combination did not record any adverse effect on litter quality and RD titer.

Keywords: β -glucan, Broiler, Immune Status, Litter Moisture, Litter pH, Mannan oligosaccharides (MOS)



Introduction

Mannan oligosaccharide (MOS) is a mannan based carbohydrate derived from the cell wall of yeast *Saccharomyces cerevisiae* (Iqbal *et al.*, 2015). It is considered that MOS exerts its effect on growth by accelerating the maturation of GIT microflora (Yang *et al.*, 2007). Dietary MOS also modulates mucin synthesis by increasing the expression of different goblet cell types and up-regulating MUC2 and mRNA expression (Chee *et al.*, 2010).

The supplementation of β -glucans protects against a number of economically-important pathogens and improves gut health and increases disease resistance (Huff *et al.*, 2010; Shao *et al.*, 2013). Yeast β -glucans supplementation enhances lysozyme and complement levels, improves phagocytic activity and alters tight junction protein expressions (Chuammitri *et al.*, 2011). Broilers supplemented with yeast β -glucans have increased humoral and cell-mediated immune responses (Kumar *et al.*, 2011; Shao *et al.*, 2013). In addition, supplementation with yeast cell wall extract prevents production loss, regulates intestinal innate immune responses (Yitbarek *et al.*, 2012). There are very few previous studies reports dealing with single and combined effects of dietary mannan- oligosaccharides (MOS) and β -glucan on immune status and litter quality in chicken. Therefore, study was planned to evaluate the effects of dietary oligosaccharides (MOS) and β -glucans on immune status and condition of litter in broiler.

Materials and Methods

The experiment was carried out at Broiler unit of Department of Poultry Science, COVAS, Parbhani. The quality feed ingredients were procured from local market and rations were prepared as per BIS (2007), at feed mixing plant COVAS, Parbhani. The experimental design and composition of various treatment diets fed to the experimental birds are given in Table 1 and 2, respectively. The data obtained was statistically analyzed by using Randomized block design (Snedecor and Cochran, 2002)

Table 1: Experimental design for housing of broiler with supplementation MOS, β -glucan and their combination

Treatment group	Treatment group details	No. of birds/pen/replication	No. of replicates	Total No. of birds
A	Control diet	20	4	80
B	Control diet + β -glucans @ 5g/100kg	20	4	80
C	Control diet + MOS@ 50g/100kg	20	4	80
D	Control diet + β -glucans @ 5g/100kg + MOS @ 50g/100kg	20	4	80
	Total	4	16	320

The pre-starter, starter and finisher rations were offered for first seven days, 8th day to 20th day and 21st to 42nd day of age, respectively. Mannan-oligosaccharide (MOS) and β -1,3-glucan were procured from B.V. BIO-CORP PVT. LTD, Pune-412206

Table 2: Percent ingredient and nutrient composition of (basal diet) pre-starter, starter and finisher rations with supplementation of MOS, β -glucan and their combination.

Feed Ingredients	Pre-Starter	Starter	Finisher
Maize	524.48	571.04	623.04
Soybean meal	402.32	372.05	310.51
Oil (veg)	31.32	17.24	28.96
Salt	3.8	3.8	3.8
DL-Methionine	2.04	2.21	1.89
Di-Calcium Phosphate	19.97	17.24	16.17
Shell grit	10.6	11.45	10.89
Trace mineral mixture ¹	1	1	1
AB2D3K ²	0.15	0.15	0.15
B-Complex ³	0.1	0.1	0.1
Choline Chloride, 60%	0.5	0.5	0.5
Toxin Binder	2	2	2

Antibiotic	0.5	0.5	0.5
L-lysine HCL	0.72	0.41	0
Cocciostat	0.5	0.5	0.5
Total	1000	1000	1000
Nutrient Composition (calculated)			
ME (kcal/kg)	2960	3060	3160
Protein (%)	23	21	19.5
Calcium (%)	0.9	0.85	0.8
Available phosphorus (%)	0.45	0.4	0.38
Lysine (%)	1.36	1.2	1.07
Methionine (%)	0.56	0.55	0.5

Temperature-Humidity Index (THI)

Average of weekly temperature humidity index were recorded in the broiler shed and tabulated as below.

Table 3: Average week wise Temperature –Humidity Index recorded in shed at 2.30 PM

THI Weeks						
	1	2	3	4	5	6
THI	80.89	80.74	83.92	83.16	83.37	87.17
THI	80.83	82.24	84.13	85.17	84.21	85.29
THI	81.42	82.18	83.99	82.97	85.14	84.8
THI	81.8	82.3	83.78	82.37	84.84	86.69
THI	80.74	83.48	84.88	85.72	85.31	91.24
THI	81.64	83.47	84.49	83.17	86.56	87.91
THI	82.11	83.88	85.48	84.29	86.67	84.71
Average	81.34	82.61	84.38	83.83	85.16	86.83

- i) Mild heat stress caused with THI: 72-79; ii) Moderate heat stress caused with THI : 80-88;
iii) Sever heat stress caused on birds with THI: 89 and above

The experiment was approved by Institutional Animal Ethics Committee vide resolution no. **IAEC/52/19 dated 2/3/2019**. The standard feeding, watering, floor space and vaccination schedule were followed.

Immune Status (RD Titre)

Serum sample from one bird per replicate was collected on the 21st and 42nd day post R.D. vaccination. Antibody titre against RD was estimated in the laboratory of Poultry Diagnostic and Research Centre of Venkateshwara Hatcheries Pvt. Ltd., Loni-Kalbhori, Pune, as per O.I.E. procedure (1992)

Moisture Percentage of Litter Material

The litter moisture was estimate as per the standard procedure AOAC, (1990) at 42nd Day of age as per the method described by Brake *et al.* (1992). Before taking the sample litter was raked. Litter samples were collected from five locations within each pen and thoroughly mixed to obtain litter material representative of the entire pen. At least 200g of litter was taken and immediately brought to the laboratory for moisture estimation. Samples were kept at 100±5°C for 12 hours in hot air oven and then cooled in a desiccator and the moisture content was determined as the weight loss during the heating-

$$\text{Moisture \%} = \frac{\text{Loss in weight } (W1g - W2g)}{\text{Weight of sample } (W1 - Wg)} \times 100$$

Where,

Wg = Weight of empty tin
 W1G = Weight of empty tin + Sample before drying
 W1-Wg = weight of the sample taken
 W2g = Weight of empty tin + Sample after drying
 W1g-W2g = Loss in weight.

Litter pH

The litter pH was measured according to the method described by Brake *et al.* (1992). The upper 10cm of litter was collected at each sample position and transported back to the laboratory for determination of pH. The pH of each litter type was measured after litter samples of 10g were suspended for 30 min in 100mL of deionized water and stirred for 5 min using pH meter. pH value was recorded until constant values were obtained (LABMAN 920 Precision pH/ OPH meter). The data collected were subjected to statistical analysis by using Randomized Block Design by Snedecor and Cochran (2002). The treatment means were compared by Critical Differences (CD) and Analysis of Variance.

Results and Discussion

Immune status (RD Titre)

The means of antibody titre against Newcastle disease (ND) at 21st and 42nd day of age were presented (Table 4). The non-significant influence ($P<0.01$) was revealed by supplementation of MOS, β -glucans and their combination on ND titre at 21st and 42nd days of age. The non-significant influences of supplementation of MOS, β -glucans and their combination on ND could be attributed to the severed heat stress caused on birds. The heat stress might have suppressed the immunity. The THI was moderate to severe in broiler shed during the experimental period (Table 3). These results are in agreement with Lopes *et al.* (2009). They concluded that non-significant influence on supplementation of β -glucans on RD antibody titer. In contrast, An *et al.* (2008) observed significantly ($P<0.05$) higher antibody titer by supplementation of β -glucans. The results in present study are not in agreement with Vesna *et al.* (2007). They concluded that the supplementation of MOS in diet significantly increased antibody titer against ND. Similar results were also revealed by Sadeghi *et al.* (2013), Waqas *et al.* (2018).

Table 4: Antibody titre of broilers against Ranikhet disease at 21st and 42nd day of age as influenced by dietary supplementation of MOS, β -glucan, and their combination

Treatments	RD Titer @ 8 HAU			
	21 st day		42 nd day	
	Mean	SE	Mean	SE
A	1	0.65	1	0.65
B	0.5	0.58	4	1
C	0	0.5	11	4
D	0	0	8	7

Table 5: ANOVA for Antibody titre of broilers against Ranikhet disease at 14th and 21st day of age as influenced by dietary supplementation with β - glucans, MOS and their combination

Sources	21st day				42nd day		
	Df	SS	MSS	F	SS	MSS	F
Treatments	3	2.75	0.917	1.571	232	77.333	0.595
Errors	12	7	0.583	-	1560	130	
Total	15		-	-			

Table 6: Moisture, percentage and pH of litter at 42nd days for different age groups supplemented with β - glucans, MOS and their combination

Moisture percentage of litter				
Samples no.	Treatment			
	A	B	C	D
I	23.25	23.31	22.51	19.60
II	21.82	24.21	23.33	19.23
III	23.64	17.76	17.63	23.37
IV	20.69	22.49	20.62	18.14
Overall mean	22.35	21.94	21.02	20.08
pH of Litter				
I	6.41	6.89	6.90	6.40
II	6.39	7.11	6.87	6.37
III	6.29	7.13	6.62	6.30
IV	6.53	7.89	6.31	7.07
Overall mean	6.40	7.25	6.67	6.78

Table 7: ANOVA for moisture percentage and pH of Litter at 42nd age groups supplemented with β - glucans, MOS and their combination

Moisture percentage				
Sources	Df	SS	MSS	F
Treatments	3	12.252	4.084	0.753
Replica	12	65.08	5.423	
Total	15	77.332	0.001	
pH of Litter				
Sources	Df	SS	MSS	F
Treatments	3	1.5092	0.503067	1.989717
Replica	12	3.034	0.252833	
Total	15	4.5432		

Moisture Percentage of Litter Materials

The supplementation of MOS, β -glucans and their combination did not influence litter moisture, however, numerically lower litter moisture was recorded for these treatment groups indicating the supplementation of MOS, β -glucans and their combination inhibited the colonization of gastrointestinal tract by pathogenic microorganism, thereby reduced viscosity of gut and ultimately resulted in low moisture percentage of litter. MOS significantly increased the goblet cell numbers. The goblet cells are responsible for the production of mucin, which binds pathogenic microorganisms and reduce their colonization of the gut mucosa (Blomberg *et al.*, 1993). The lower litter moisture obtained in β -glucans, MOS and their combination groups might be due to their effect of on the mucosal layer of the intestine. The present findings are in agreement with Sarica and Cam (1998) who reported non-significant findings with the litter moisture, Stanley *et al.* (2004) reported non-significant differences in the litter moisture when diet was supplemented with yeast cell culture, Cheraghi *et al.* (2014) and Yalcin *et al.* (2014) also reported non-significant findings with the litter moisture when the diet was supplemented with yeast cell wall / yeast extracted β -glucans. However, Rezaei *et al.* (2011) reported that the inclusion of micronized insoluble fibre @ 0.5 percent resulted in significant ($p < 0.05$) decrease in the litter moisture. Hassan and Ryu (2013) revealed that fecal ammonia and CO₂ gas emission was significantly ($P < 0.05$) decreased in multi-prebiotics diet supplemented group. Sadeghi *et al.* (2013) reported that inclusion of prebiotic-based mannan-oligosaccharide and β -glucan resulted in

lowest litter moisture percent compared with other treatment groups. However, Onwurah *et al.* (2013) revealed that supplementation of yeast in diet was resulted to void more water in their droppings may be due to more heterotrophic gastrointestinal tract of the birds fed yeast content diet.

Litter pH

The supplementation of MOS, β -glucans and their combination did not influenced pH values. However, a pH value of litter was within limit of ideal conditions. The findings of the present study are in agreement with, Meluzzi *et al.* (2008) who reported that the litter pH was unaffected by both rearing condition and litter type. Miles *et al.* (2011) concluded that lower NH_3 production in the litter is indicative of ideal litter pH. Cheraghi *et al.* (2014) reported non-significant findings with litter pH when the diets were supplemented with yeast extracted β -glucans and α -mannans. Tasistro *et al.* (2007) indicated that Sodium bisulphate can be a best alternative to suppress the NH_3 emission from the litter, Ritz *et al.* (2009) stated that the litter moisture should not exceed 25 percent as excess litter moisture may increase ammonia emission (>25 ppm) from the litter and thereby the pH of litter may also be altered. Youssef *et al.* (2011) revealed that ammonia was produced in wet litter because of microbial activity on uric acid and thereby increased the pH of litter .

Conclusion

It may be concluded that the supplementation of β -glucans, MOS and their combination did not record any adverse effect on litter quality as well as litter pH. However, there were significant differences ($P<0.01$) between antibody titer against Ranikhet disease at 21st day of age.

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Conflict of Interests

There is no conflict of interest.

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