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SINGLE AND COMBINED EFFECTS OF MANNAN OLIGOSACCHARIDE (MOS) AND DIETARY PROTEIN ON THE PERFORMANCE AND IMMUNITY RESPONSE OF LAYING HENS

By

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Abstract: This experiment was conducted to study the single and combined effects of mannan oligosaccharide and dietary protein on the performance, egg quality, immunity response, serum constituents and economical efficiency of laving hens which designed as factorial arrangement, three levels of crude protein (CP: 14.75,13.25, and 11.75%) x two levels of mannan oligosaccharide (MOS: 0 and 0.1%). A total number of 72 Hy-Line W- 36 laying hens 49 weeks old were distributed randomly into six equal groups each group containing 12 hens. The results obtained could be summarized as follows: 1- Level of CP% significantly (P<0.01) affected each of egg production % (EP%), egg mass (EM), egg weight (EW), feed conversion (FC) and caloric conversion ratio (CCR). The level of 14.75% CP had numerically higher EP%, EM, EW than the level 13.25% CP and it had better (P < 0.01) FC or CCR than the other levels of CP%. The level of 13.25% CP caused insignificant increase in immune responses of SRBCs, CBH response of laying hens. 2-Adding MOS to layer's diets significantly improved FC, crude protein conversion (CPC) and CCR. Also, it caused significant increases in shell %, yolk%, yolk index (YI), serum protein and albumin while decreased egg albumen%. total MOS supplementation caused insignificant increase in primary and secondary immune response of SRBCs and CBH response of hens compared with unsupplemented diet. 3-Hens fed the 14.75% CP unsupplemented diet with MOS had significantly higher (P < 0.01) EP% and EM than the groups fed 11.25% CP with or without 0.1% MOS diets, but, it insignificantly differed than other groups. The group of 14.75% CP + 0.1% MOS had the highest (P < 0.01) EW, the lowest FI(P < 0.05) and the best (P < 0.01) FC and CCR followed by group 13.25% CP + 0.1% MOS of EW, FC and CCR. 4-Eggs produced by layers fed diet contains 13.25% CP+0.1% MOS had significantly higher shell% but lower albumen% than other groups. Eggs produced by group11.75% CP+0.1% MOS had the higher YI than other

groups. 5-The highest values of primary immune response against sheep red blood cells (SRBCs) were found in the groups fed diets contained 13.25% CP with or without MOS, whereas the highest value of secondary immune response was found in the group fed diet contains 11.75% CP+ 0.1% MOS followed by 13.25% CP+0.1% MOS group which had the highest value of cutaneous basophil hypersensitivity (CBH) response. The highest values (P < 0.01) of serum total protein and albumin were of groups 14.75 % CP+0.1% MOS and 13.25% CP+0.1% MOS. These results favored the middle levels of CP (13.25%) + 0.1% MOS in diet of laving hens caused better effect in immunity responses of hens than others and the highest values of economical efficiency with the relative economical efficiencies were shown in the groups 14.75 % CP+0.1% MOS and 13.25% CP+0.1% MOS. It could be concluded that adding 0.1% MOS as natural feed additive in diets of laying hens which contain a level of CP that recommended by strain catalog or less with 1.5% were economically better in production and *immunity without adverse effect on egg quality.*

INTRODUCTION

Using alternatives to antibiotic growth promotants in commercial chickens have become important mainly because of apprehensions about the possible development of resistant bacteria. At the same time, continuous use of antibiotic growth promotants in breeders may have one important ramification that could affect the poultry industry, reduction in the efficacy of antibiotics when used in progeny that are hatched to the same parents Devegowda, (Shashidhara and 2003). Also, antibiotic growth promotants resulted in the occurrence of resistant microorganisms which became one of the major problems in human medicine. One of many such classes of alternatives is the prebiotic mannan oligosaccharide (MOS). The MOS is derived from the outer cell wall of yeast. Mannose, the main component of MOS, is a unique sugar because many enteric bacteria have receptors that bind to it (Griggs and Jacob, 2005). The MOS supplementation is considered because it is not only shifts gastrointestinal microflora balance toward beneficial organisms (Spring et al., 2000, Fairchild et al., 2001) but also resulted in significant improvement in antibody responses in broiler and layers (MacDonald, 1995, Savage et al., 1996, Cotter, 1997, Cotter et al., 2000 Cotter et al., 2002, Raju and Devegowda, 2002). Supplementation of poultry diets with MOS results in improved production in terms of body weight gain and feed conversion (Parks et al., 2001), which partly may be due to its hypothesized nutrient sparing effect and primarily to its influence on nutrient utilization in the

gastrointestinal tract (Kumprecht et al., 1997, Savage et al., 1997, Sonmez and Eren, 1999).

Dietary protein content has a much consideration due to its high cost and its great effect on the production parameters of laying hens. Matching dietary protein and amino acids levels to the production requirement of laying hens is another important mean of reducing nitrogen emissions and excretion. Lowering the CP of the laying hens diets not only reduce nitrogen consumption but also means that less unutilized nitrogen is excreted. The response by the laying hens to dietary protein levels has been controversial for many years. Fernandez, et al. (1973) reported that increasing dietary protein level lead to an increase in egg production %. Also, average egg weight of layers increased as dietary protein level increased (Summers, 1993). Moreover, Calderon and Jensen (1990) observed an improvement in FC due to high dietary protein level. However, Angelovicova (1994) found that a low-protein diet containing 14.1 % CP reduced average daily FI and improved FC. Glick, et al. (1983) showed that diet deficient in protein (33% of requirement) could reduce numbers of lymphocytes in the thymus of chickens. However, the responses were varied by strain, dietary protein (Cheema et al., 2003) environment, stress, production state and health status. Thus, protective immune responses require a supply of nutrients at the appropriate times and amounts (Humphrey et al., 2002). The present experiment aimed to study single and combined effects of mannan oligosaccharide and dietary protein on the performance and immunity responses, serum constituents, and economical efficiency of laying hens

MATERIALS AND METHODS

The experimental work of the present study was carried out at the Poultry Research Station, Poultry Department, Faculty of Agriculture. Fayoum University from April to July 2003, to study single and combined effects of mannan oligosaccharide and dietary protein on the performance, egg quality, immunity response, serum constituents of laying hens. A total number of 72 Hy- Line W- 36 laying hens at 49 weeks old were distributed randomly into six equal groups each group containing 12 hens in 12 replicates, one hen / replicate. The experiment was designed as factorial arrangement, three levels of CP (14.75, 13.25, and 11.75%) x two levels of mannan oligosaccharide (0 and 0.1%). The basal diets were formulated to satisfy nutrient requirements of laying hens according to the strain catalog recommendations (14.7 CP % and 2770 ME. K cal / kg). The composition and chemical analyses of the experimental diets are shown in Table 1. Artificial light was used beside the normal day light to provide 16-hour day photoperiod. Feed and water were provided *ad libitum*. Individual body

weights were recorded at the beginning and the end (61 weeks of age) of the experiment to calculate live body weight changes (LBWC). Egg number (EN) and egg weight (EW) were recorded daily to calculate egg production % (EP% = EN*100/84 day) and egg mass (EM= EN* EW). Feed intake (FI) was recorded weekly and it using to calculate feed conversion (FC= FI/EM), crude protein conversion (CPC= FI* CP%/ EM) and caloric conversion ratio (CCR= FI* ME. K cal / EM).

Egg quality measurements were determined monthly on eggs of the last three days. Twelve eggs / group were collected monthly throughout the experimental periods to determine egg shape index % (SI, Carter, 1968), shell thickness (ST) including shell membranes was measured using a micrometer at three locations on the egg (air cell, equator and sharp end), the percentage of shell, albumen and yolk were calculated .Yolk color (YC) was determined by matching the yolk with one of the 15 bands of the "1961- Roche improved yolk color fan. Yolk index % was calculated according to (Well, 1968), Haugh unit score (HU) was applied from a special chart using egg weight and albumen height which was measured by using a micrometer according to Haugh (1937).

At 54 weeks of age four hens of each group were injected in wing vein with 0.2 ml of sheep red blood cells solution (SRBCs 9% suspension) and the blood samples were collected from the wing vein of these birds after one week to determine SRBCs primary immune response. The same birds were reinjected at 60 weeks of age and the blood samples were collected from these birds after 5 days to determine SRBCs secondary immune response in serum and determine the serum constituents. Antibody response against SRBCs were measured in serum using micro haemagglutination technique as described by **Yamamoto and Glick (1982) and Dix and Taylor (1996).** The titers were expressed as the log 2 of the reciprocal of the highest dilution giving visible agglutination (**Atta et al., 1998).** Serum constituents were determined calorimetrically using kits according to **Weichselbaum (1946)** for total protein, **Dumas and Biggs (1972)** for albumin **, Allain, et al. (1974)** for total cholesterol and **Wahlefeld (1974)** for triglycerides.

To determine cutaneous basophil hypersensitivity (CBH) response, three hens from each group were randomly selected at 61 weeks of age and injected with 0.1 ml of phytohaemagglutinin –P (PHA-P,100 μ g/ml) subcutaneously in the right toe web, whereas. 0.1 ml saline was injected subcutaneously in the left toe web which served as the control. The thickness of both toe webs were measured in mm using a micrometer at 24 hr after injection. The CBH response was calculated as described by (Atta *et al.*, 1998) as follows:

CBH response = Thickness of right toe web (PHA-P) response / Thickness of left toe web (saline response)

Economical efficiency of egg production was calculated from the inputoutput analysis which was calculated according to the price of the experimental diets and eggs produced. These values were calculated as the net revenue per unit of total cost. Analysis of variance was computed using the general linear model (GLM) procedure of statistical analysis system according to **SPSS (1999)**. Significant differences among means were evaluated using Duncan's multiple range test (**Duncan, 1955**).

RESULTS AND DISCUSSION

Productive performance of laying hens

Results presented in Table 2 showed that levels of CP% significantly (P<0.01) affected each of EP%, EM, EW, FC and CCR. The level of 14.75% CP had insignificantly (P>0.05) higher EP%, EM, EW than the level of 13.25% CP and, it had better FC or CCR than the other levels of CP%. However, the 11.75% CP had the lowest EP%, EM, EW (56.00%, 2413g and 51.42g respectively) associated with the worst FC and CCR being 3.13 g feed/ g EM and 8.68 calorie/ g EM. No significant effects of dietary protein on the FI, CPC and LBWC of laying hens were found. Hammershoj and Kjaer (1999) and Bunchasak et al. (2005) reported trend that increasing the egg weight with increasing dietary protein however, decreasing the level of protein improved CPC . Similar trends were found by Bunchasak et al. (2005) that high CP of 16 and 18 % tend to have better EP% and EM than the lower level of 14 % CP .However, Hammershoj and Kjaer (1999) reported that different levels of dietary protein did not affect EP%. On the other hand, Harms and Russell (1995) reported that the 10.95% CP satisfied the requirements needed for egg production, egg mass and egg content.

Adding MOS to diets of laying hens significantly improved FC, CPC and CCR than the unsupplemented diet (2.54 g feed/g EM, 0.342 g CP/g EM and 7.06 calorie/ g EM vs 2.86 g feed/g EM, 0.376 g CP/g EM and 7.94 calorie/ g EM respectively), regardless of level CP level. Whereas, the MOS supplementation had no significant effects on EP%, EM, EW, FI and LBWC. Similarly, **Shashidhara and Devegowda (2003)** found that MOS supplementation had no influence on egg production of broiler breeders. On the contrary, **Guerrero ,1995, Berry and Lui , 2000 and Stanley** *et al.,2000* reported considerable improvement in egg production in the MOS-fed birds. Similar results of significant FC improvement were found in laying hens and Japanese quail at the same level of MOS supplementation

were reported by Chukwu and Stanley (1997) and Ghosh, *et al.* (2007). This is may be because MOS maintain gut health by adsorption of pathogenic bacteria containing type-1 fimbrae of different bacterial strains and remove the bacteria from gut (Oyofo *et al.*, 1989 and Spring *et al.*, 2000) and increase villus height, uniformity and integrity also, increase in crypt depth is attributed to greater expenditure of energy to develop the absorptive surface (Dawson and Tricarico, 2002, Loddi *et al.*, 2002, Olivera, *et al.*, 2006, Mourao *et al.*, 2006 and Ghosh, *et al.*, 2007)

Except for LBWC and CPC, there were significant treatment interaction effects on each of EP%, EM, EW, FI, FC, and CCR as shown in Table 2. The 14.75% CP unsupplemented diet with MOS had significantly higher EP% and EM (70.83% and 3118g) than either the 11.25% CP unsupplemented diet with MOS (54.17% ad 2380g) or the 11.75% CP supplemented 0.1% MOS (57.84% and 2445g) whereas the former insignificantly differed than other groups for these traits. The level of 14.75% CP supplemented 0.1% MOS had significantly higher EW (55.46), however the level of 11.75% CP supplemented 0.1% MOS had lower EW (50.37g) than other groups. The level 14.75% CP unsupplemented MOS consumed more daily FI (94.12) than other groups whereas the 14.75% CP supplemented 0.1% MOS had the lowest FI of 85.24g. Therefore, the latter group had the best (P<0.01) FC of 2.17g feed/ g EM followed by 13.25% CP+0.1% MOS being 2.48 g feed/ g EM however poorer FC than other groups was shown by the 11.75% CP unsupplemented MOS being 3.28 g feed/ g EM. Similar trend were found for CCR.

Egg quality

Except YI, levels of CP% insignificantly affected all external and internal egg quality traits. The 11.75% CP had the highest YI of 56.82% as shown in Table 3. However, Hammershoj and Kjaer (1999) reported that with increasing dietary protein, albumen quality traits and egg shell % decreased. MOS supplementation resulted in significant increases in shell %, yolk% and YI of 10.73, 29.49 and 56.71%, respectively and a decrease in albumen% being 59.81% (Table 3). Similarly, Berry and Lui (2000) and Shashidhara and Devegowda (2003) reported that the MOS improved egg shell quality traits in older breeder females. There were significant differences in each of shell%, albumen% and YI (P<0.05) due to treatment interactions, may be due to improvement in calcium availability. Eggs produced by layers fed the 13.25% CP+0.1% MOS had significantly higher shell% but lower albumen% than other groups (11.27% and 59.46%). Eggs produced by hens fed the11.75% CP+0.1% MOS had the highest YI of 59.88% whereas these fed 13.25%CP unsupplementation the

MOS produced significantly lower YI of 50.49% than other groups as shown in Table 3.

Immunity responses

Results presented in Table 4 indicated that the levels of CP% significantly affected the primary immunity. The level of 13.25% CP caused an increase in primary immune response ($P \le 0.071$) of SRBCs of laying hens, however, insignificantly influenced either the secondary immune response of SRBCs or the value of CBH response of laying hens. MOS supplementation caused insignificant increase in primary ($P \le 0.227$), secondary (P≤0.055) immune response of SRBCs and CBH response (P≤0.126) of hens, regardless of CP%. However, Malzone et al. (2000) and Shashidhara and Devegowda (2003) reported that hens fed diet supplemented with 0.05% MOS had higher SRBCs titers than controls at one week post-sensitization and significant increase in the values of antibody titers of broiler breeders. This effect on antibody titers may be due to the influence of the MOS on immune system and/or improvement of intestinal absorption of some related nutrients, such as Zn, Cu, Se. In addition to the reduction of the pathogenic bacteria load in the intestine and preventing the acute immune response against such bacteria (Finucane, et al., 1999 and Spring et al., 2000). The interaction between levels of CP% and levels of MOS% showed no consistent trend in the values of primary and secondary immune response of SRBCs and CBH response. The highest values of primary immune response were found in the groups which fed diet contains 13.25% CP with or without MOS, whereas the highest value of secondary immune response was found in the group fed the 11.75% CP+ 0.1% MOS diet followed by the group fed diet contains 13.25% CP+0.1% MOS. The highest value of CBH response was found of the group fed diet contains 13.25% CP+0.1% MOS. The results indicate to the middle levels of CP (13.25%) + 0.1% MOS in diets of laying hens caused better effect in immunity responses of hens than others.

Serum constituents

The levels of CP % significantly affected triglycerides (P \leq 0.05), the lowest value was found in the group fed the 13.25% CP. However, there were no effect of the different levels of CP% used in this study on each of serum total protein, albumin, globulin, A/G ratio and cholesterol. Adding of 0.1% MOS in diets of laying hens caused significant increases (P<0.01) in serum total protein and albumin compared with the hens fed the unsupplemented diet (10.09 vs 8.98g/dl and 6.36 vs 5.16g/dl), regardless of CP % however, insignificantly affected other serum constituents as shown

in Table 4. Significant differences were found (P<0.01) in serum total protein and albumin due to levels of CP% and MOS % interaction, the highest values were of groups 14.75 % CP+0.1% MOS and 13.25% CP+0.1% MOS being 10.34 and 10.24g/dl for total protein and 6.54 and 6.67 24g/dl for serum albumin, whereas, the lowest values were 8.68g/dl and 4.51g/dl of total protein and albumin for group 11.75 % CP+0.0MOS, respectively as shown in Table 4. it can be concluded that adding MOS to diets containing different levels of CP may be induced such improvement in synthesis of serum total protein and albumin.

Economical efficiency

Supplementing layer diets which differed in level of CP with 0.1% MOS improved economical efficiency and relative economical efficiency of laying hens compared with those fed the unsupplemented diet. The highest values of economical efficiency of 1.515 and 1.365 with the relative economical efficiencies of 116.00 and 104.49 were shown for the groups fed 14.75% CP+0.1% MOS and 13.25% CP+0.1% MOS. However, the worst values were found (0.967 and 74.01 of economical efficiency and its relative economical efficiency) for the group fed 11.75% CP+0.0 MOS as shown in Table 5. Similarly, **Namra (2006)** suggested that incorporation of 0.15 % baker's yeast in diet of quail layers apparently exhibited better amelioration in feed cost /egg than the control group. It could be concluded that adding 0.1% MOS as natural feed additive in diets of laying hens which contain a level of CP that recommended by strain catalog or less with 1.5% were economically better in production and immunity without adverse effect on egg quality.

Ingredients	14.75% CP	13.25% CP	11.75 % CP
Yellow corn, ground	69.30	71.42	73.47
Soybean meal (44%CP)	20.00	15.34	10.44
Wheat bran	0.00	2.36	5.06
Calcium carbonate	8.00	8.10	8.10
Di calcium phosphate	2.00	2.00	2.00
Vit. and Min. premix*	0.30	0.30	0.30
Sodium chloride	0.30	0.30	0.30
DL- methionine	0.10	0.12	0.15
Lysine	0.00	0.06	0.18
Total	100.00	100.00	100.00
Calculated analysis**			
CP %	14.69	13.19	11.63
EE%	2.79	2.91	3.03
CF%	2.92	2.90	2.90
Ca%	3.43	3.46	3.45
Available P %	0.47	0.47	0.45
Methionine%	0.35	0.34	0.35
Methionine+Cystine%	0.61	0.57	0.55
Lysine%	0.72	0.67	0.68
ME, K cal /kg	2767	2765	2760
Cost (L.E./ton)***	816.3	793.0	779.4
Relative cost****	100.00	97.15	95.48

Table (1): Composition and chemical analyses of the experimental diets.

* Each 3.0 Kg of the Vit. and Min. premix manufactured by Agri-Vet Company, Egypt contains : Vit. A, 10000000 IU Vit. D₃ 2000000 IU Vit. E, 10 g Vit. K₃, 1 g Vit. B1, 1 g Vit. B2, 5 g Vit. B6, 1.5 g Vit. B12, 10 mg choline chloride, 250 g biotin, 50 mg folic acid, 1 g nicotinic acid, 30 g Ca pantothenate, 10 g Zn, 50 g Cu, 4 g Fe, 30 g Co, 100 mg Se, 100 mg I, 300 mg Mn, 60 g, and completed to 3.0 Kg by calcium carbonate. *** According to NRC, 1994. **** According to market prices of 2003.

**** Assuming that the control equals 100.

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C		(()		
		Shell						
Items	Shape index	thickness	Shell %	Albumen%	Yolk %	yolk color	yolk index	Hough unit
		(mm)						
Level of CP %			-					
14.75	76.73±0.42	$0.352{\pm}0.004$	10.41 ± 0.18	60.53±0.35	29.22 ± 0.32	$9.69 {\pm} 0.13$	$53.30{\pm}0.92$ ^B	79.35±1.59
13.25	76.19 ± 0.42	$0.358 {\pm} 0.004$	10.79 ± 0.18	59.99±0.35	29.22±0.32	$9.89 {\pm} 0.13$	52.50±0.92 ^B	76.23±1.58
11.75	76.99±0.42	0.351 ± 0.004	$10.24{\pm}0.18$	$60.89 {\pm} 0.35$	28.71±0.32	9.61±0.13	56.82±0.92 ^A	81.02 ± 1.58
Level of MOS %	-		-		-			
0.00	76.92±0.34	$0.355 {\pm} 0.003$	10.24 ± 0.14 ^b	61.10±0.27 ^a	28.60 ± 0.25 ^b	$9.68 {\pm} 0.11$	51.89 ± 0.68 ^B	79.41±1.31
0.10	76.92±0.34	$0.353 {\pm} 0.003$	10.73 ± 0.14 ^a	59.81 ± 0.27^{b}	29.49±0.25 ^a	$9.84{\pm}0.11$	56.71±0.68 ^A	78.33±1.30
Interaction								
14.75% CP+0.0 MOS	77.07±0.59	$0.357 {\pm} 0.01$	10.17 ± 0.24^{b}	$61.02{\pm}0.46^{\mathrm{ab}}$	28.71±0.45	9.52 ± 0.18	$51.49{\pm}1.02^{\rm ed}$	81.66±2.27
13.25% CP+0.0 MOS	77.04±0.59	$0.355 {\pm} 0.01$	10.31 ± 0.24^{b}	$60.52{\pm}0.46^{\rm abc}$	29.08 ± 0.44	$10.04 {\pm} 0.18$	$50.49{\pm}1.02^{d}$	77.39 ± 2.23
11.75% CP+0.0 MOS	76.66±0.59	$0.352{\pm}0.01$	$10.22\pm0.24^{\mathrm{b}}$	61.78±0.46 ^a	28.03±0.44	$\textbf{9.48} \pm \textbf{0.18}$	$53.76{\pm}1.02^{\rm bc}$	79.26±2.23
14.75% CP+0.1%MOS	76.40±0.59	$0.347{\pm}0.01$	$10.30{\pm}0.24^{\rm b}$	$60.04{\pm}0.46^{ m bc}$	29.72±0.44	$\textbf{9.85} \pm \textbf{0.18}$	56.07 ± 1.02^{b}	77.14±2.23
13.25% CP+0.1%MOS	75.34±0.59	0.361 ± 0.01	$11.27{\pm}0.24^{\rm a}$	59.46±0.46°	29.36±0.44	9.93 ± 0.18	54.52 ± 1.02^{b}	75.06 ± 2.23
11.75% CP+0.1%MOS	77.32±0.59	0.351 ± 0.01	$10.60{\pm}0.24^{\rm ab}$	$60.01 \pm 0.46^{ m bc}$	29.40 ± 0.44	9.74 ± 0.18	59.88±1.02ª	82.79±2.23
Over all mean	76.64±0.24	$0.354{\pm}0.002$	$10.48{\pm}0.10$	60.46 ± 0.19	$29.05{\pm}0.18$	9.76 ± 0.08	54.30 ± 0.46	$78.87 {\pm} 0.91$

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a.c and A. C values in the sam	Ver all mean 683+	11.75% CP+0.1%MOS 7.00±0	13.25% CP+0.1%MOS 8.00±0	14.75% CP+0.1%MOS 7.00±0	(1.75% CP+0.0 MOS 6.50±((3.25% CP+0.0 MOS 8.00±([4.75% CP+0.0 MOS 4.50±(Interaction).10 7.33±().00 6.33±(Level of MOS %	(1.75 6.75±((3.25 8.00±((4.75 5.75±(Level of CP %	imm	Ttoma Prin	Table (4): Single and combined
1.3/ 8.4	1 27 84). 90 10.). 90 9.0). 90 8.5). 90 7.5). 90 8.5	0.90 7.0).57 9.1	0.57 7.6).65 8.7).65 8.7).65 7.7		unity i	nary S	effects of ma
± ∪.00 hin the same i	2+0 66	00 ± 0.94	0±0.94	0±0.94	0±0.94	0±0.94	0±0.94		7±0.52	7 ± 0.52		5±0.69	5±0.69	5±0.69		mmunity	econdary	nnan oligosac
1.120±0.02 tem followed by	1 126+0 02	1.003 ± 0.05	1.291 ± 0.05	$1.201{\pm}~0.05$	$1.115 {\pm} 0.05$	$1.065 {\pm} 0.05$	$1.080 {\pm} 0.05$		$1.165 {\pm} 0.04$	$1.087 {\pm} 0.04$		$1.059 {\pm} 0.05$	$1.178 {\pm} 0.05$	$1.140 {\pm} 0.05$		response	CBH	charide and die
y.so±v.iv different superscri	0 56±0 10	$9.83{\pm}0.24^{\mathrm{AB}}$	$10.24{\pm}0.23^{\rm A}$	$10.34{\pm}0.26^{\rm A}$	$8.68 \pm 0.23^{\circ}$	$8.94 \pm 0.23^{\circ}$	9.33±0.23 ^{вс}		$10.09 {\pm} 0.14^{ m A}$	$8.98{\pm}0.14^{\rm B}$		9.32±0.20	9.59±0.22	9.76±0.23		Protein g/dl	Total	etary protein on ir
5./0±0.20	5 76+0 26	6.01±0.44 ^a	6.67±0.48 ^a	6.54±0.54 ^a	4.51±0.48°	5.36 ± 0.48^{ab}	5.59±0.48 ^{ab}		$6.36{\pm}0.28^{\mathrm{A}}$	$5.16{\pm}0.28^{\rm B}$		5.34 ± 0.33	6.02 ± 0.36	6.00 ± 0.39		(A) g/dl	Albumin	nmune respons
3./8±0.15	3 78+0 15	3.82 ± 0.33	3.57±0.37	$3.80{\pm}0.42$	4.17±0.37	3.58±0.37	3.73±0.37		3.73 ± 0.30	3.83 ± 0.30		3.98 ± 0.35	3.57±0.37	3.76 ± 0.39		(G) g/dl	Globulin	se and serum co
1.82±0.18 t P <0.05 for a 1	1 87+0 18	2.00 ± 0.38	2.40±0. 43	1.76 ± 0.49	1.14±0. 43	2.02±0. 43	1.53 ± 0.43		$2.07{\pm}0.24$	1.56 ± 0.24		1.61 ± 0.28	2.21 ± 0.30	1.63 ± 0.32		ratio	A/G	onstituents of la
235.20±18.49	735 70+18 40	231.86 ± 40.06	222.97±44.79	244.19±51.72	226.45±44.79	209.30±44.79	276.45±44.79		231.97±24.23	237.40±24.23		229.46±27.86	216.13±29.54	262.62±31.59		mg/dl	Cholesterol	ying hens (Mean.
573.83±4.97	303 83+4 07	393.90 ± 10.77	369.51±12.04	392.07±13.90	415.24±12.04	376.52±12.04	415.24±12.04		385.32±7.77	402.34±7.77		403.39±9.21 ^a	373.02±8.62 ^b	405.31±8.12 ^a		mg/dl	Triglycerides	±SE).

0.070	1.000	0.000			0.720	$\frac{1}{2} = \frac{1}{2} = \frac{1}{2}$
6.598	7.960	8.366	5.591	7.532	8.425	Net revenue / hen (L.E.) $f - c = g$
12.145	13.792	13.887	11.375	13.750	14.875	Total price of eggs / hen (L.E.) d x $e = f$
0.25	0.25	0.25	0.25	0.25	0.25	Price /egg (L.E.) e
48.58	55.17	55.55	45.50	55.00	59.50	Total number of eggs /hen d
0.010	0.000	0.700	U.ULU	0.210	0.707	$I = \frac{1}{2} $
ハ ハフタ	7 805	887 2	7 817	6 7 1 8	6 454	Total fand cost than $(I \in I)$ as $h = c$
6.856	7.087	6.527	7.425	7.841	7.905	Total feed intake hen(kg) b
0.809	0.823	0.846	0.779	0.793	0.816	Price/ k feed(LE) a
+0.1% MOS	+0.1% MOS	+0.1% MOS	+0.0% MOS	+0.0% MOS	+0.0% MOS	
11.75%CP	13.25%CP	14.75%CP	11.75%CP	13.25 %CP	14.75%CP	Items
m±SE).	otein diets (Mea	charide in low pr	mannan oligosaco	effect of adding	aying hens under the	Table (5): Economical efficiency of la

e.....(according to the local market price at the experimental time).

g/c(net revenue per unit feed cost).

r.....(assuming that economical efficiency of control group).

REFERENCES

- Allain, C.C., Poon, L. S., Chan, C. S. G., Richmond, W., Fu, P. C. (1974). Enzymatic determination of total serum cholesterol. Clin. Chem., 20: 470-475.
- Angelovicova, M. (1994). Economic use of a low-protein feed mixture in layer diet. Zivocisna-Vyroba., 39: 1049 1062.
- Atta, A.M. Abdou, A.M. Mohamed, F.R. and Goher, N.E. (1998). Immunological variation among commercial broiler strains. Egypt. J. Anim. Prod., 35: 113-124.
- Berry,W. D. and Lui, P. (2000). Egg production, egg shell quality and bone parameters in broiler breeder hens receiving Bio- Mos and Eggshell 49. Poult. Sci., 79 (Suppl. 1):124. (Abst.).
- Bunchasak, C., Poosuwan, K. and Nukraew, R. (2005). Effect of Dietary protein on egg production and immunity responses of laying hens during peak production period. Int. J. Poult. Sci., 4: 701-708.
- Calderon, V. and Jensen, L.S. (1990). The requirement for sulfur amino acid by laying hens as influenced by the protein concentration. Poult. Sci., 69: 934-944.
- Carter, T. C. (1968). The hen egg. A mathematical model with three parameters. Br. Poult. Sci., 9: 165 171.
- Chukwu, H. I. and Stanley, V. G. (1997). Dietary Saccharomyces cervisiae and mannan oligosaccharide reduced the deleterious effect of heat stress on White Leghorn laying hens. Association of research Directors Eleventh Biennial Research Symposium October 1-4, 1997.
- Cheema, M. A., Qureshi, M.A. and Havenstein, G.B. (2003). A comparison of the immune profile of commercial broiler strains when raised on marginal and high protein diets. Int. J. Poult. Sci., 5: 300-312.
- Cotter, P. F. (1997). Modulation of the immune response: Current perceptions and future prospects with an example from poultry. pp. 195–204 in Biotechnology in the Feed Industry. T. P. Lyons and K.A. Jacques, ed. Nottingham University Press, Nottingham.
- Cotter, P. F., Malzone, A., Paluch, B., Lilburn, M. S. and Sefton. A. E. (2000). Modulation of humoral immunity in commercial laying hens by a dietary prebiotic. *Poult. Sci.*, 79(Suppl. 1):38 (Abst.).

Laying, crude protein, Mannan oligosaccharide, immune, serum constituents

- Cotter, P. F., Sefton, A. E., and Lilburn. M.S.(2002). Manipulating the immune system of layers and breeders: Novel applications for mannan oligosaccharides. pp. 21–28 in Nutritional Biotechnology in the Feed and food Industries.
- **Dawson K. A. and Tricarico J. (2002).** The evolution of yeast cultures 20 years of research. Navigating from Niche Markets to Mainstream Proceeding from Alltech's 16th Annual European, Middle Eastern and African Lecture Tour.
- Dix, M.C. and Taylor, R. L. J. (1996). Differential antibody responses in 6.8 maior histocompatbility .B complex congenic chickens. Poult. Sci., 75: 203 207.
- Dumas, B.T. and Biggs, H.G. (1972). In Standard Methods of Clinical Chemistry. Vol 7. Academic Press New York. USA.
- Duncan, D.B. (1955). Multiple range and multiple F Test. Biometrics, 11: 1-42.
- Fairchild, A. S., Grimes, J. L., Jones, F. T., Wineland, M. J., Edens, F. W., and Sefton. A. E. (2001). Effects of hen age, Bio-Mos, and Flavomycin on poult susceptibility to oral Escherichia coli challenge. Poult. Sci., 80:562–571.
- Fernandez, R. Salman, A. J. and Ginnis, J.M. (1973). Effect feeding different protein levels and of changing protein level on egg production. Poult. Sci., 52: 64-69.
- Finucane, M., Spring, P. and Newman, E. (1999). Incidence of mannose sensitive adhesions in enteric bacteria. Poult. Sci., 78 (Suppl.): 139 (Abst.).
- Glick, B. Taylor, R.L. J. Martin, D.E. Watabe, M. Day, E.J. and Thompson, D. (1983). Caloric-protein deficiencies and immune response of the chicken. II. Cell mediated immunity. Poult. Sci., 62: 1889 – 1893.
- Ghosh, H. K., Halder, G. Samanta, G., Paul, S. K. and Pyne, S. K. (2007). Effect of dietary supplementation of organic acid and mannan oligosaccharide on the performance and gut health of Japanese Quail (Coturnix Coturnix Japonica). Asian J. Poult. Sci., 1: 1-7.
- Griggs, J. P. and Jacob J. P.(2005). Alternatives to antibiotics for organic poultry production. J. Appl. Poult. Res., 14: 750-756.
- Guerrero, M. R. (1995). Using yeast culture and lactic acid bacteria in broiler breeder diets. pp. 371–378 in Biotechnology in the Feed

Industry. T. P. Lyons and K. A. Jacques, ed. Nottingham University Press, Nottingham.

- Hammershoj, M. and Kjaer, J. B. (1999). Phase feeding for laying hens: Effect of protein and essential amino acids on egg quality and production. Acta Agriculturae Scandinavica, Section A- Animal Sciences., 49: 31-41.
- Harms, R. H. and Russell, G. B. (1995). A re-evaluation of the protein and lysine requirement for broiler breeder hens. *Poult. Sci.* 74: 581-585.
- Haugh, R. R. (1937). The Haugh unit for measuring egg quality. US Egg Poult. Mag., 43: 552-555.
- Humphrey, B.D. Koutsos, E.A. and Klasing, K.C. (2002). Requirements and priorities of the immune system for nutrients Nutrition biotechnology in the feed and food industries. Proceedings of Alltech's 18th annual symposium pp. 69 – 77 Lyons T.P. Jasques K.A. Nottingham. UK Nottingham University Press.
- Kumprecht, I., Zobac, P., Siske, V., Sefton, A. E and Spring, P. (1997). Effects of dietary mannanoligosaccharide level on performance and nutrient utilization of broilers. Poult. Sci. 76 (Suppl. 1):132. (Abstr.).
- Loddi, M. M., Nakaghi, I. S. O., Edens, F., Tucci, F. M., Hannas, M. I., Moraes, V. M. B. and Ariki, J. A. (2002). Effect of mannan oligosaccharides and organic acids on intestinal morphology integrity of broiler evaluated by scanning electron microscope. Proceeding of 11th European Poultry Science Conference, Bremen, Germany, PP: 121.
- MacDonald, F. (1995). Use of immunostimulants in agricultural applications. pp. 97–103 in Biotechnology in the Feed Industry. T. P. Lyons and K.A. Jacques, ed. Nottingham University Press, Nottingham.
- Malzone, A., Paluch, B., Lilburn, M.S. and Sefton, A. E. (2000). Modulation of humoral immunity in commercial laying hens by dietary probiotic. Poult. Sci., 79 (suppl 1), 165.
- Mourao, J.I., Pinheiro, V., Alves, A., Guedes, C.M., Pinto, L., Saavedra, M.J., Spring, P., and Kocher, A. (2006). Effect of mannan oligosaccharides on the performance, intestinal morphology and cecal fermentation of fattening rabbits. Anim. F. Sci. and Tech., 126: 107-120.

- Namra, M. M. (2006). Influence of using Baker,s Yeast and microbial phytase in Japanese Quail diets on productive performance and some physiological parameters. Egypt. Poult. Sci., 26: 579 607.
- Oliveira, M. C., Gravena, R. A., Marques, R. H., Cancherini, L. C., Rodrigues, E. A. (2006). Morphometry of intestinal mucosa in 21 day- old broiler chickens fed mannan-oligosaccharides and a blend of enzyme. Poultry Science Association 95th Annual Meeting Abstract., July 16-19, 2006.
- Oyofo, B. A., Deloach, J. R., Corrier, D.E., Norman, J. O. Ziprin, R. I. and Mollenhauer, H. H. (1989). Prevention of Salmonella typhimurium colonization of broilers with D-mannose. Poult. Sci., 68:1357–1360.
- Parks, C. W., Grimes, J. L., Ferket, P. R. and Fairchild, A. S. (2001). The effect of mannan oligosaccharides, bambermycins, and virginiamycin on performance of large white male market turkeys. Poult. Sci., 80:718–723.
- Raju, M. V. I. N. and Devegowda, G. (2002). Esterified-Gluco-mannan in broiler chickens diets-contamimated with aflatoxin, ochratoxin and T-2 toxin: Evaluation of its binding ability (in vitro) and efficacy as immunomodulator. Asian-Aust. J. Ani. Sci., 15: 1051-1056.
- Savage, T. F., Cotter, P. F. and E. I. Zakrzewska (1996). The effect of feeding mannanoligosaccharide on immunoglobulins, plasma IgG and bile IgA of wrolstadMWmale turkeys. Poult. Sci., 75(Suppl. 1):143. (Abst.).
- Savage, T. F., Zakrzewska, E. I. and J. R. Andreasen (1997). The effects of feeding mannan oligosaccharide supplemented diets to poults on performance and morphology of small intestine. Poult. Sci., 76(Suppl. 1):139. (Abst.).
- Shashidhara, R. G. and Devegowda, G. (2003). Effect of dietary mannanoligosaccharide on broiler breeder production traits and immunity. Poult. Sci., 82:1319.
- Sonmez, G. and Eren, M. (1999). Effects of supplementation of zinc bacitracin, mannanoligosaccharide and probiotic into the broiler feed on morphology of the small intestine. Vet. Fak. Derg. Uludag Univ., 18:125–138.
- Spring, P., Wenk, C., Dawson, K. A. and Newman, K. E. (2000). The effects of dietary mannan oligosaccharides on cecal parameters and

the concentrations of enteric bacteria in the ceca of Salmonellachallenged broiler chicks. **Poult. Sci., 79:205–211.**

- SPSS (1999). Statistical software package for the social sciences SPSS, Int., USA.
- Stanley, V. G., Brown, C. and Sefton, T. (2000). Single and combined effects of dietary protease and mannanoligosaccharide on the performance of laying hens. Poult. Sci., 79(Suppl. 1):62. (Abstr.).
- Summers, J.D. (1993). Reducing excretion of the laying hen by feeding lower crude protein diets. Poult. Sci., 72: 1473 1478.
- Wahlefeld, A.W. (1974). In Methods of Enzymatic Analysis. Vol 5. HU Bergmever. Ed. Academic Press. New York., 1831-1835.
- Weichselbaum, T.E. (1946). An accurate and rapid method for the determination of protein in small amounts of blood serum and serum. Am. J.Clin. Path., 16: 40-48.
- Well. R. J. (1968). The measurement of certain egg quality: A study of the hens egg. Ed. By T.C. Carter Pub. Oliver and Boy Edinbrugh pp. 220-226 and 235-236.
- Yamamoto, Y. and Glick, B. (1982). A comparison of the immune response between two lines of chickens selected for differences in the weight of the bursa of fabricius. Poult. Sci., 61: 2129 - 2132.

التأثيرات الفردية و المشتركة للمنان اوليجوسكريد و بروتين العليقة

على الأداء والإستجابة المناعية للدجاج البياض

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أجريت هذه التجربة لدراسة التأثيرات الفردية و المشتركة للمنان اوليجوسكريد وبروتين العليقة علي الأداء والإستجابة المناعية للدجاج البياض وصممت كتوزيع عاملي، ثلاث مستويات من البروتين الخام (14.75، 13.25، 11.75%) x مستويين من المنان اوليجوسكريد (صفر ،0.1%). 72 دجاجة بياضة عمر 49 أسبوع من سلالة الهاي لين دبليو 36 وزعت عشوائياً إلى 6 مجاميع متساوية كل مجموعة احتوت على 12 دجاجة.

ويمكن تلخيص النتائج كما يلى

 1-مستوى البروتين كان له تأثير معنوي جدا على كلا من نسبة إنتاج البيض ، كتلة البيض، وزن البيض ، معدل التحويل الغذائي ومعدل تحويل الطاقة. المستوى 14.75 % بروتين هو الأعلى رقميا في نسبة إنتاج البيض ، كتلة البيض، وزن البيض مقارنة بالمستوى 13.25 % بروتين و المستوى 14.75 % بروتين هو الأحسن (معنويا جدا) في معدل التحويل الغذائي ومعدل تحويل الطاقة ذلك مقارنة بالمستويات الأخرى من الهروتين والمستوى 13.25 % بروتين سبب زيادة

غير معنوية في الإستجابات المناعية ضد كر ات الدم الحمر اء للغنم و استجابة فرط الحساسية للدجاج البياض .2- إضافة المنان اوليجوسكريد لعلائق الدجاج البياض حسنت معنويا كلا من معدل التحويل الغذائي ، معدل تحويل البروتين و معدل تحويل الطاقة ايضا سبب زيادة معنوية في نسبة قشرة البيض ، نسبة الصفار ، دليل الصفار ، البروتين الكلى بالسيرم والبيومين السيرم وسبب نقص في البيومين البيضة. إضافة المنان اوليجوسكريد سبب زيادة غير معنوية في الإستجابات المناعية الأولية والثانوية ضد كرات الدم الحمراء للغنم و استجابة فرط الحساسيةً للدّجاجات البياض مقارنة بعدم الإضافة. 3- الدجاجات المغذاة على عليقة بها 14.75 %بروتين وغير مضاف لها المنان اوليجوسكريد كانت الأعلى معنويا جدا في نسبة إنتاج البيض ، كتلة البيض مقارنة بالمجمو عات المغذاة على علائق بها 11.75 % بروتين ومضاف أو غير مضاف لها 0.1 % منان اوليجوسكريد ولكن لم تختلف معنويا عن المجموعات الأخرى. مجموعة 14.75 % بروتين المضاف لها 0.1% منان اوليجوسكريد هي الأعلى (معنويا جدا) في وزن البيضة و الأقل (معنويا) في كمية العلف المأخوذ والأحسن (معنويا جدا) في معدل التحويل الغذائي ومعدل تحويل الطاقة يليها مجموعة ال 13.25 % بروتين + 0.1% منان اوليجوسكريد في وزن البيضة ، معدل التحويل الغذائي ومعدل تحويل الطاقة. 4- البيض المنتج من دجاجات مغذاة على عليقة محتوية على 13.25 % بروتين + 0.1% منان اوليجوسكريد كان أعلى معنويا في نسبة القشرة ولكن الأقل في نسبة الألبيومين عن المجموعات الأخرى. والبيض المنتج من مجموعة 13.25 % بروتين +0.1% منان اوليجوسكريد كان أعلى معنويا في دليل الصفار. 5- القيم الأعلى للإستجابة المناعية الأولية ضد كرات الدم الحمراء للغنم وجدت في المجموعات المغذاة على علائق بها 13.25 % بروتين مضاف أو غير مضاف لها منان اوليجوسكريد في حين أن القيمة الأعلى للإستجابة المناعية الثانوية وجدت في المجموعة المغذاة على عليقة بها 11.75 % بروتين +0.1 % منان اوليجو سكريد يلها مجموعة ال 13.25 % بروتين + 0.1% منان اوليجوسكريد والتي امتلكت أعلى قيمة لإختبار فرط الحساسية. القيم الأعلى (معنويا جدا) للبروتين الكلي بالسيرم والبيومين السيرم كانت للمجمو عات 14.75 % بروتين +0.1% منان اوليجوسكريد و 13.25 % بروتين +0.1% منان اوليجوسكريد. هذه النتائج ايدت ان المستوى الأوسط من البروتين (13.25 %) +0.1 % منان اوليجوسكريد في عليقة الدجاج البياض سبب أحسن تأثير في الإستجابات المناعية للدجاجات عن المجمو عات الأخرى . والقيم الأعلى للكفاءة الإقتصادية ونسبة الكفاءة الإقتصادية كانت في المجموعتين % 14.75 بروتين +0.1% منان اوليجوسكريد و 13.25 % بروتين +0.1% منان اوليجوسكريد.

بذلك يمكن التوصية بإضافة 0.1% منان اوليجوسكريد كإضافة غذائية طبيعية في علائق الدجاج البياض والتي تحتوي على مستوى البروتين الموصى به تبعا لكتالوج السلالة أو أقل منة بمقدار 1.5% حيث كانوا الأحسن اقتصاديا في الإنتاج ومناعيا بدون تأثير سلبي على جودة البيضة.

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