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Immunomodulating effect of B-glucans and mannan oligosaccharide on broiler chicks vaccinated with Newcastle disease virus

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This study was carried out to determine the immunomodulating effect of β -glucans and mannan oligosaccharide (MOS) on the immune response of chickens to Newcastle disease vaccine. The results showed that birds received β -glucans and MOS having higher average body weights values and significantly higher ND HI antibody titer than the other non medicated groups. Thymus, spleen and bursal indices of control negative showed significantly lower values than vaccinated medicated and non-medicated groups. Both total and differential leukocytic and lymphocytic counts showed significantly higher in medicated group than other groups. Liver function test showed lower AST and ALT in medicated group than other groups. Results of challenge test with NDV confirmed that MOS and B glucans immunostimulant improved protection rate by 15% in medicated than non- medicated ones. In conclusion MOS and B glucans can be given to chicken to improve both body weight and protection against VV NDV challenge that predominated in Egypt.

Commercial poultry flocks receive a lot number of vaccines to protect them from environmental pathogens; therefore, a great effort had been expanded to develop strategies to enhance chicken immune response, especially in facing immunosuppression caused by extraneous agents, infections, intoxication or by certain vaccine viruses. Immunomodulation could improve vaccinal immunity and possibly selectively promote responses that are critical for protection.

Immunomodulators usually classified according to their origin into biological and chemical products (Poli, 1984). This classification further broken down into physiological products, substances of microbial origin and synthesis compounds.

The mannan-oligosaccharide (MOS) is derived from the outer cell wall of yeast, and its evaluation in diets for breeders is of particular interest because it not only shifts gastrointestinal microflora balance toward beneficial organisms (Spring *et al.*, 2000; Fairchild *et al.*, 2001) but also has immunomodulatory properties (Cotter *et al.*, 2002). The yeast cell wall has powerful antigenic stimulating properties, and it is well established that this property is a characteristic of the mannan chain (Ballou, 1970). This study was carried out to determine the immunostimulant effects of commercial feed additive preparations containing a mannanoligosaccharide plus β -glucans on chicken, performance and immune response to ND vaccine. Body weight gain, HI and challenge With NDV that endemic in Egyptian poultery farms as well as bursal, thymic and spleen body weight ratio were taken as criteria for evaluation based.

Materials and methods

Immunostimulants (ALPHAMUNE®). It's a commercial feed additive product composed of (1-3, 1-6) β -glucans and (MOS) obtained from Alpharma Animal Health. USA (patch NO AG51242).It was used in ration at a rate of 500 gram/ ton of finished fed.

Chickes. A total number of 225 one day-old commercial (white HI-line rooster) chickens obtained from El-Wady Company were used in this study.

Newcastle disease (ND) vaccinal strains. 1-Hitchiner B1 and La Sota strains, produced by Pfizer International Company, USA with each vial contain virus titre of 10^9 EID_{50} was used after titration for vaccination of experimental chicks via eye instillation route.

Clone 30. Vaccine nobilis clone 30 (Lot No: 06829AJ01, Intervet international B.V. Boxmeer – Holland) with virus titer of 10^6 EID_{50} was used for vaccination of experimental chicks via eye instillation

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Velogenic NDVs. A local velogenic viscerotropic Newcastle disease virus (vvNDV) isolate (Shible and Reda, 1976) was kindly supplied by Newcastle Diseases Department; Veterinary Serum & Vaccine Research Institute, Abbasia, Cairo. Egypt were used for challenge test.

Haemagglutination (HA) and haemagglutination inhibition (HI) tests. HA and HI test were carried out according to (Anon. 1971).

Haemtological studies. Total leukocytic count was counted according to the technique described by (Nutt and Herrick, 1952) while differential leukocytic count was done by the standard method of Battelnent described (Schalm, 1973).

Biochemical analysis. ALT and AST were done according to the method of (Reitman and Framkel, 1957). Serum uric acid was done according to the method descried by (Barham and Trinder, 1972) and serum creatinin was done according to the method described by (Houot, 1985).

Bursa body weight index. It was calculated according to (Ying *et al.*, 2003) as following: Bursa: body weight ratio = bursa weight/ body weight. Bursal index = Bursa: body weight ratio X 1000.

Challenge test. Chickens were challenged via intramuscular route. Each chicken received a dose of 0.5ml / bird containing $10^6 \text{ EID}_{50 \text{ VV NDV}}$ according to (Afify, 1990). Birds with persisted symptoms till the end of the observation period were considered as if dead.

Statistical analysis. Statistical analysis of variance (ANOVA) test was used to estimate differences among treatments according to (Steel and Torrie, 1960).

Experimental design. The used chicks (225) were floor reared and fed on balanced commercial ration free from antimicrobial agents. At the 1st day of life, 5 chicks were sacrificed for organ body weight ratio and serum while the rest of obtained chicks were divided into 4 groups (1-4). Groups (1 and 3) containing 60 chicks each, while groups 2 and 4 containing 50 chicks each. Each group was kept in a separate clean disinfected room.

Chicks of groups 1 and 3 were feed on ration without additives, while those of groups 2 and 4 were feed on ration supplemented with the immunostimulant (Alphamune®), in dose of 0.5 gm/ kg. At the 5th day of age all chicks were S/C

vaccinated with inactivated avian influenza (H5N1) (0.3 ml/ bird). At the 7th day of age, chicks of group (1 and 2) were kept ND nonvaccinated control while birds of groups (3 and 4) were vaccinated each with 10^6 EID₅₀ Hitchiner B1 via eye instillation and revaccinated at the 18th day of age as each bird was given 10^6 EID₅₀ La Sota via eye instillation route. At 35 day of age 20 birds from groups were separated and challenged with 0.5 ml containing 10⁶ VVND. Challenged chickens were kept under daily observation for 21 days with daily record of symptoms, deaths and post - mortem lesions. Ten birds from group 1 and 3 were left without challenge to be control.

Experimental chicken groups were weekly subjected to the following: Life body weight of random 5 birds / group as well as weight of bursal, thymus and spleen of each bird was recorded to calculate organ body weight ratio. Random 5 non-coagulated blood samples on EDTA were collected for total anddifferential leukocytic count. Random coagulated 5 blood samples from wing vein were collected for serum collection. The collected serum samples were divided into two equal quantities, labeled and stored at-20°C until use. The collected sera were tested for detection of NDV HI as well as liver and kidney function test.

Results and Discussion

numbers of There are large immunostimulatory components were reported to be used for stimulating the chicken immune response to face the problem of vaccination failure, which constitute a challenge to poultry industry all over the world. The application of immunostimulant is not only to raise resistance of birds but also to improve their immune response to vaccination (Afify, 1990; Awaad et al., 2000). The work was designed to evaluate the effect of Alphamune as immunomodulator in chickens, where data presented in (Table 1) showed that administration of Alphamune was significantly increased body weight at 7 days old 80.30±1.88 gm verses 72.90 ±1.58 for control group. While from 14 to 35 days there is no significant difference could be detected values of different groups as well as that of control group. This result agrees with those of Solis de los (Santos et al., 2007) where a significant weight difference at 7 days only between treated and non treated poults was found while no difference at 3 weeks old.

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Group	Tre	atment			Ag	e / week		
No.	I.S	Vacc	0	1	2	3	4	5
1	-	-	36.39±1.2	72.90±1.58	130.00 ± 2.31	221.80±8.40	319.30±4.29	428.40±9.93
2	+	-		80.30±1.88*	133.00±3.21	228.50 ± 5.5	332.00±5.09	446.25±17.95
3	-	+			135.40 ± 2.85	225.00±4.18	325.60±5.26	428.60±15.93
4	+	+			137.60±2.35	229.60±5.28	343.50±12.04	467.50±14.16

Table (1): Effect of immunostimulant on average body weight of ND vaccinated and non-vaccinated chickens.

Each value represents mean \pm S.E.

* Significant difference between groups by t-student test at $P \le 0.05$.

Table (2): The effect of immunostimulant on mean ND HI antibody titer in vaccinated and non-vaccinated chickens.

Group	Tre	atment			А	ge / week		
No.	I.S	Vacc	0	1	2	3	4	5
1	-	-	7.8±86	6 ± 0.32	$#4 \pm 0.55 \text{ b}$	# 2.4±0.51b	# 1.6±0.51b	# 1.6±0.51b
2	+	-		6.2±0.37	$4.4 \pm 0.51b$	$2.8 \pm 0.37b$	$1.8 \pm 0.37b$	$1.8 \pm 0.55b$
3	-	+		6 ± 0.32	6.4± 0.51a	$7.2 \pm 0.58a$	6.6± 0.69a	$5.6 \pm 04a$
4	+	+		6.2±0.37	7.2±0.37a	7.8± 0.58a	$7.2 \pm 0.37a$	$6.2 \pm 0.49a$

Each value represents mean \pm S.E.

#: Significant variation between groups by ANOVA test at $P \le 0.05$.

Different superscript letters a and b denote significant variation respectively by LSD at $P \le 0.05$.

 Table (3): Effect of immunostimulant on thymus index of vaccinated and non-vaccinated chickens with NDV live vaccine.

Group	Trea	tment			Ag	e / week		
No.	I.S	Vacc.	0	1	2	3	4	5
1	-	-	4.60±0.12	5.60 ± 0.06	#6.00±0.10b	#6.10±0.08b	#6.70±0.12c	#5.60±0.18c
2	+	-		5.9±0.07*	6.60±0.09a	7.21±0.15a	7.65±0.23b	5.80±0.20b
3	-	+			6.63±.0.11a	7.25±0.20a	7.91±0.22ab	6.33±0.21ab
4	+	+			6.71±0.10a	7.50±0.2a	8.35±0.30a	6.50±0.25a

Each value represents mean \pm S.E.

* Significant difference between groups by t-student test at $P \le 0.05$.

#: Significant variation between groups by ANOVA test at $P \le 0.05$.

Different superscript letters a, b and c denote significant variation respectively by LSD at $P \le 0.05$.

Table (4): Effect of immunostimulant on mean spleen index of vaccinated and non-vaccinated chickens with NDV live vaccine.

Group	Trea	tment			Age	/ week		
No.	I.S	Vacc.	0	1	2	3	4	5
1	-	-	0.50 ± 0.01	1.00 ± 0.05	#1.50±0.08c	#1.90±0.10c	#2.05±0.08c	2.30±0.05
2	+	-		1.35±0.07*	1.70±0.07c	2.10±0.10bc	2.10±0.15c	2.34 ± 0.06
3	-	+			2.30±0.11b	2.46±0.12b	2.50±0.11b	2.40 ± 0.09
4	+	+			2.63±0.15a	2.87±0.17a	2.90±0.18a	2.44±0.15

Each value represents mean \pm S.E.

* Significant difference between groups by t-student test at $P \le 0.05$.

#: Significant variation between groups by ANOVA test at $P \le 0.05$.

Different superscript letters a, b and c denote significant variation respectively by LSD at P \leq 0.05.

Table (5): Effect of immunostimulant on mean bursal index of vaccinated and non-vaccinated chickens with NDV live vaccine.

Group	Trea	tment			Age	e / week		
No.	I.S	Vacc.	0	1	2	3	4	5
1	-	-	1.80 ± 0.08	2.50±0.10	#2.80±0.15c	#3.20±0.12b	#2.41±0.15c	#1.55±0.11b
2	+	-		2.85±0.12*	2.90±0.14c	3.40±0.12b	2.60±0.13bc	1.85±0.12b
3	-	+			3.20±0.13b	3.80±0.14ab	3.00±0.19b	2.50±0.15a
4	+	+			3.43±0.13a	4.10±0.18a	3.55±0.21a	2.75±0.20a

Each value represents mean \pm S.E.

* Significant difference between groups by t-student test at $P \le 0.05$.

#: Significant variation between groups by ANOVA test at $P \le 0.05$.

Different superscript letters a, b and c denote significant variation respectively by LSD at $P \le 0.05$.

Group	Treat	tment			Age	/ week		
No.	I.S	Vacc.	0	1	2	3	4	5
1	-	-	16.1±1.30	14.3±1.10	#11.6±0.50b	#12.1±0.74b	#13.1±0.51b	#13.2±0.9b
2	+	-		22.5±1.30**	17.3±1.30ab	15.6±0.75ab	14.8±0.8ab	13.4±0.95b
3	-	+			20.6±1.15a	21.5±1.20a	22.2±1.23a	21.1±1.25b
4	+	+			26.2±1.20a	28.5±1.9a	32.3±1.7a	30.6±2.3a

Table (6): Effect of immunostimulant on mean total leucocytic count X 10³ of vaccinated and non-vaccinated chickens with NDV live vaccine.

Each value represents mean \pm S.E.

** Significant difference between groups by t-student test at $P \le 0.01$.

#: Significant variation between groups by ANOVA test at $P \le 0.05$.

Different superscript letters a, b and c denote significant variation respectively by LSD at $P \le 0.05$.

Regarding thymus index, spleen index and bursal index of chicken fed on Alphamune supplemented ration and vaccinated with NDV revaled significant increase in values than results of other groups where it give 6.50 ± 0.25 , $2.44 \pm$ 0.15 and 2.75 \pm 0.20 respectively at 35 days of age (Table 3-5). These results come in agreement with the finding of Ying et al., (2003) where mean percentage of organ body weight ratios of liver, spleen, Kidney, thymus and bursa of Fabricius exhibited a significant (P<0.05) increase in MOS as compared to those of control group. Results of total and differential leucocytic count (Table 6, 7) were significantly higher TLC on group 4 at 35 days of age $(30.6 \pm$ 2.3) in comparison to $(13.2 \pm 0.9, 13.4 \pm 0.95,$ 21.1 ± 1.5) for the other groups. It was observed that the source of increased in TLC is the significantly increased lymphocyte counts due to use of Alphamune®, Increased TLC in group 4 can be attributed to immunostimulation effect of compounds of Alphamune. This result was previously observed by (Fleischer et al., 2000; Acevedo et al., 2001) who recoded increased TLC with administration of MOS and b-glucan respectively. Chicken group 4 that fed on Alphamune® supplemented ration and vaccinated with NDV vaccine showed significant lower AST and ALT levels at 35 days of age (Table 8, 9). Where the results is $172.17 \pm$ 7.15 and 9.85 \pm 0.20 respectively verses 201.56 \pm 7.53 and 14.3 \pm 0.25 in untreated vaccinated group .This result was observed by (Santhosh et al., 2003) in treated group with MOS all over the breeding period.

Statistical analysis of uric acid and creatinine values (Table 10, 11) resulted in non significant difference between different groups up to 35 days of age. The instability in creatinine value from week to week may be related to change in feed and protein concentration. Birds of group (4) showed significant HI titers to NDV at 35 days of age than other groups (Table 2); this higher HI titers resulted in 95% protection in this group 4 compared to 85% protection in group 3, 20% protection in group 2 and 0% protection in group 1 (Table 13). Our results clearly showed the specific immune stimulation and protection against challenge in group 4 were attributed to B-glucan compound of Alphamune due to increasing functional activity of macrophage and neutrophils. Yun *et al.*, (2003); Sakurai *et al.*, (1992) reported that orally B-glucan indirectly stimulate the immunity in the respiratory system of mice by activating macrophage in the payer's patches of the gut.

From the above discussed data we could conclude that Alphamune[®] could increase body weight gain, improve immunity of the birds and decrease susceptibility to NDV challenge.

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						5	roup No. 5	Group No. and treatment	nt						
Group 1: Control	Control			Group 2: I.S	I.S			Group 3: vaccinated	accinated			Group 4: I.	Group 4: I.s +vaccination	0 U	
week H	L	Μ	E	Н	Г	Μ	E	Η	Γ	Μ	E	Н	Г	Μ	E
20.5	74.7	2.9	1.9	20.5	74.7	2.9	1.9	20.5	74.7	2.9	1.9	20.5	74.7	2.9	1.9
$25.8^{\pm1.08}$	67.8 ± 1.07	3.6 ± 0.24	2.8 ± 0.2	17.2 ± 0.58	77.8*±0.37	3±0.32	2±0.32	25.8*±1.08	67.8±1.07	3.6 ± 0.24	2.8 ± 0.2	17.2 ± 0.58	77.8*±0.37	3 ± 0.32	2 ± 0.32
23.8 ± 1.02	73.4±0.75	2.0±0.32	1.2 ± 0.37	$20.4{\pm}0.51$	75.6*±0.4	2.2±0.37	1.8 ± 0.2	$45.8^{\pm0.37}$	48.0 ± 0.89	$3.4^{*\pm0.24}$	2.8*±0.37	$41.6^{\pm 0.51}$	52.1±0.32	2.7±0.37	$3.6^{\pm0.4}$
20.4 ± 0.81	76.4*±0.51	2.6±0.24	1.2 ± 0.58	18.2 ± 0.37	77.2*±0.37	3.4±0.6	0.6 ± 0.24	$49.8^{\pm 0.37}$	43.6±0.24	2.8 ± 0.37	4.0*±0.2	$44.8^{\pm}0.37$	49.6 ± 0.24	2.2±0.37	3.4*±0.24
17.6 ± 0.51	76.8*±0.37	2.8±0.51	2.8±0.37	18.4 ± 0.51	77.6*±0.51	1.8 ± 0.2	2.2±0.37	46.6*±0.75	48.6±0.51	2.2 ± 0.37	2.6±0.24	$34.1^{*}\pm0.32$	59.5*±0.52	2.2±0.37	2.6 ± 0.24
20.4 ± 0.93	74.4±0.93	1.8 ± 0.2	2.4±0.24	17.2±0.86	77.6*±1.12	2.4±0.37	2.8±0.37	42.0*±0.71	52.2±0.58	1.8 ± 0.2	3.6±0.37	33.6*±0.57	61.0*±0.71	1.8 ± 0.2	3.4±0.4
nificant diffe Heterophils	erence at P<	0.05 betwi nphocytes	een treatec	d and non ti onocytes;	reated group E = Eosino	s. phils.									
le (8): Effec	ct of immui	nostimula	ant onme	an of AST	in sera of v	vaccinate	d and noi	n-vaccinate	ed chicken	is with NE	JV live va	accine.			
;	Treatme	ent						Ag	e / week						
ond no. –	I.S V	acc.	0		1		2			3		4		S	
1	ı	1	$101.30\pm$	4.15	131.30±6.	21	#155.00±	⊧5.90 a	#162.3	0±5.35 a	#17	74.55±5.84 b		<u>90.10±6.8</u> 2	2 a
7	+	ı			125.32 ± 5.3	85	$130.10 \pm$	6.15 b	142.60)±5.60 b	159	9.21±6.17 b		58.32±6.25	p
e	ı	+					$160.30 \pm$	7.51 a	171.25	5±5.85 a	192	4.80±8.13 a		01.56±7.53	а
4	+	+					$136.50 \pm$	6.22 b	139.50)±5.51 b	162	2.23±6.54 b		2.17±7.15	ab
value repres gnificant var erent supersc	sents mean	-S.E. en groups ı,b and c d	by ANOV enote sign	VA test at F ufficant vari	≥≤ 0.05. iation respec	tively by I	LSD at P≤	≤ 0.05.							
le (9): Effec	ct of immur	nostimula	unt on me	an of AL7	l in vaccine	tted and n	10n-vacci	inated chic	kens with	NDV live	vaccine.				
Group No.	Tre	atment							Age / wee	k					
	I.S	Vact		0	-			2		3		4		S	
1		1	4.92	2±0.15	7.90±0.2	00	#10.1	7±0.20 a	#12.7	77±0.22 a	#1.	3.12±0.25 b		3.50±0.26	q
0	+	'			7.76±	0.21	8.65=	±0.22 b	11.2	0±0.25 b	10).57±0.22 c	12	2.30±0.26	c
ę	·	+					10.5()±0.21a	12.8	9±0.26 a	14	4.05±0.27 a	14	4.30±0.25	a
	3 20.4 ± 0.81 4 17.6 ± 0.51 5 20.4 ± 0.93 Significant diffe A = Heterophils able (8): Effect Broup No. - Group No. - Group No. Croup No. 1 1 2 3 3 3 3 4 4 4 5 Croup No. 1 1 2 2 3 3 3 3 4 4 4 5 Croup No. 1 2 3 3 3 3 5 Croup No. - 1 3 3 Croup No. - 1 1 2 2 3 3 Croup No. - 1 1 2 2 Croup No. - 1 2 2 Croup No. - 1 2 2 Croup No. - 1 2 2 Croup No. - 1 2 2 3 3 Croup No. - 1 2 2 Croup No. - 1 2 2 Croup No. - 1 2 2 3 3 Croup No. - 1 2 2 Croup No. - 1 2 2 2 2 3 3 3 2 2 2 2 2 2 2 2 2 2 2 2	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	3 20.4 ± 0.81 $76.4^{*\pm}0.51$ 26 ± 0.24 4 17.6 ± 0.51 $76.8^{*\pm}0.37$ 2.8 ± 0.51 *Significant difference at P 20.4 ± 0.93 1.8 ± 0.2 *Significant difference at P 0.05 betw *H = Heterophils; $L = Lymphocytes$ *H = Heterophils; $L = Lymphocytes$ *Table (8): Effect of immunostimula Group No. Treatment 1 - - 2 + + 4 + + #: Significant variation between groups Different superscript letters a,b and c d Different superscript letters a,b and c d I.S Vacc. 1 - - - 2 + + + + #: Significant variation between groups Different superscript letters a,b and c d I.S Vac #: 3 - + - - - - 3 - I.S I.S - - - 2 + - - - - - - -	20.4\pm0.81 76.4 ± 0.51 76.4 ± 0.51 2.6 ± 0.24 1.2 ± 0.58 17.6 ± 0.51 $76.8^{\pm}\pm0.37$ 2.8 ± 0.37 2.8 ± 0.37 20.4 ± 0.93 74.4 ± 0.93 1.8 ± 0.2 2.8 ± 0.37 20.4 ± 0.93 74.4 ± 0.93 1.8 ± 0.2 2.8 ± 0.37 20.4 ± 0.93 74.4 ± 0.93 1.8 ± 0.2 2.8 ± 0.37 $1fiftcant difference at P \leq 0.05 between treated Pin M_{0} M = M_{1} Heterophils; L = Lymphocytes; M = M_{1} e(8): Effect of immunostimulant onmea M = M_{1} M_{1} 0 up No. Ireatment 0 0 1 101.30\pm 1 101.30\pm 1 101.30\pm 2 + + + 3 + + 4 + + 1 101.30\pm 2 + + + 3 + + $	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	3 20.4±0.81 76.4*±0.51 2.6±0.24 1.2±0.58 18.2±0.37 77.2*±0.37 4 17.6±0.51 76.8*±0.51 76.8*±0.51 2.8±0.51 2.8±0.51 2.8±0.51 77.6*±0.51 76.8*±1.12 5 20.4±0.93 74.4±0.93 1.8±0.2 2.4±0.24 17.2±0.86 77.6*±0.51 \times Significant difference at P≤ 0.05 between treated and non treated group \ast H = Heterophils; L = Lymphocytes; M = Monocytes; E = Eosinon Table (8): Effect of immunostimulant onmean of AST in sera of v Table (8): Effect of immunostimulant onmean of AST in sera of v Table (8): Effect of immunostimulant onmean of AST in sera of v Table (8): Effect of immunostimulant onmean of AST in sera of v Table (8): Effect of immunostimulant onmean of AST in sera of v Table (8): Effect of immunostimulant onmean of AST in sera of v Table (8): Effect of immunostimulant onmean of AST in sera of v Table (8): Effect of immunostimulant onmean of AST in sera of v Table (8): Effect of immunostimulant onmean of AST in sera of v Table (9): Effect of immunostimulant onmean of AST in sera of v b sector of the test sub and c denote significant variation respector Table (9): Effect of immunostimulant on mean of ALT in vaccina Table (9): Effect of immunostimulant on mean of ALT in vaccina Table (9): Effect of immunostimulant on mean of ALT in vaccina Table (9): Effect of immunostimulant on mean of ALT in vaccina Table (9): Effect of immunostimulant on mean of ALT in vaccina Table (9): Effect of immunostimulant on mean of ALT in vaccina Table (9): Effect of immunostimulant on mean of ALT in vaccina Table (9): Effect of immunostimulant on mean of ALT in vaccina T = 1	3 20.4±0.81 76.4*±0.51 2.6±0.24 1.2±0.58 18.2±0.37 77.2*±0.37 3.4±0.6 4 17.6±0.51 76.8*±0.37 2.8±0.37 2.8±0.37 18.4±0.51 1.8±0.2 5 20.4±0.93 74.4±0.93 1.8±0.2 2.4±0.54 17.2±0.86 77.6*±1.12 2.4±0.37 *FI = Heterophils; L = Lymphocytes; M = Monocytes; E = Eosinophils. *H = Heterophils; L = Lymphocytes; M = Monocytes; E = Eosinophils. Table (8): Effect of immunostimulant onmean of AST in sera of vaccinate Group No. Treatment 1 101.30±4.15 131.30±6.21 1 101.30±4.15 131.30±6.21 2 + + + 4 + + Each value represents mean ±S.E. #: Significant variation between groups by ANOVA test at P≤0.05. Different superscript letters a,b and c denote significant variation respectively by 1 Table (9): Effect of immunostimulant on mean of ALT in vaccinated and r Group No. Treatment #: Significant variation between groups by ANOVA test at P≤0.05. Different superscript letters a,b and c denote significant variation respectively by 1 Table (9): Effect of immunostimulant on mean of ALT in vaccinated and r Group No. Treatment Group No. Treatment J. 4.92±0.15 7.90±0.20 1 4.92±0.15 7.90±0.20 1 4.92±0.15 7.90±0.20 1 4.92±0.15 7.90±0.20 2 +	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	3 20.4±0.81 76.4±0.51 2.6±0.24 1.2±0.58 18.2±0.37 71.2±0.37 3.4±0.6 0.6±0.24 49.8±0.37 4 17.6±0.51 76.8±0.37 2.8±0.37 2.8±0.37 2.8±0.37 46.8±0.37 5.8±0.37 46.8±0.37 5.8±0.37 46.8±0.37 5.8±0.37 45.8±0.37 45.8±0.37 2.8±0.37 45.8±0.37 45.8±0.37 45.8±0.37 45.8±0.37 45.8±0.37 45.8±0.37 45.8±0.37 45.8±0.37 45.8±0.37 42.0*±0.71 Fsignificant difference at P≤ 0.05 between treated and non treated groups. F Heterophils; L = Lymphocytes; M = Monocytes; E = Eosinophils. Fable (8): Effect of immunostimulant onmean of AST in sera of vaccinated and non-vaccinate J	$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	37 3.4 ± 0.6 0.6 ± 0.24 49.8 ± 0.37 43.6 ± 0.51 2.2 ± 0.37 $4.0^{\pm0.0}$ 51 1.8 ± 0.2 2.2 ± 0.37 46.6 ± 0.75 48.6 ± 0.51 2.2 ± 0.37 2.6 ± 0.3 51 1.8 ± 0.2 2.2 ± 0.37 42.0 ± 0.71 52.2 ± 0.58 1.8 ± 0.2 3.6 ± 0.3 roups. roups. roups. roups. roups. $\frac{1}{2}$ $\frac{2}{2.4\pm0.37}$ $\frac{2}{2.8\pm0.37}$ $\frac{4.0.6\pm0.51}{2.2\pm0.58}$ $\frac{2.8\pm0.37}{1.8\pm0.2}$ $\frac{4.0^{\pm0.6}}{3.6\pm0.2}$ $\frac{1}{3.6\pm0.3}$ $\frac{1}{3.6\pm0.5}$ $\frac{1}{3.2\pm0.58}$ $\frac{1}{3.8\pm0.2}$ $\frac{3.6\pm0.3}{3.6\pm0.3}$ $\frac{1}{3.6\pm0.3}$ $\frac{2}{3.6\pm0.5}$ $\frac{1}{3.2\pm0.58}$ $\frac{1}{3.2\pm0.5}$ $\frac{1}{3.2\pm5.58}$ $\frac{1}{3.0.10\pm6.15}$ $\frac{1}{12.56\pm5.50}$ $\frac{1}{130.50\pm5.51}$ $\frac{1}{12.2\pm0.25}$ $\frac{1}{10.17\pm0.22}$ $\frac{1}{10.50\pm0.25}$ $\frac{1}{10.50\pm0.22}$ $\frac{1}{10.50\pm0.22}$ $\frac{1}{10.50\pm0.25}$ $\frac{1}{10.50\pm0.25}$ $\frac{1}{10.50\pm0.25}$ $\frac{1}{10.50\pm0.25}$ $\frac{1}{10.50\pm0.25}$ $\frac{1}{10.50\pm0.22}$ $\frac{1}{10.50\pm0.22}$ $\frac{1}{10.50\pm0.22}$ $\frac{1}{10.50\pm0.22}$ $\frac{1}{10.50\pm0.22}$ $\frac{1}{10.50\pm0.22}$ $\frac{1}{10.50\pm0.25}$ $\frac{1}{10.50\pm0.25}$ $\frac{1}{10.50\pm0.25}$ $\frac{1}{10.50\pm0.22}$ $\frac{1}{10.50\pm0.20}$ $\frac{1}{10.50\pm0.20}$ $\frac{1}{10.50\pm0.20}$ $\frac{1}{10.50\pm0.20}$ $\frac{1}{10.50\pm0.20}$ $\frac{1}{10.50\pm0.20}$ $\frac{1}{10.50\pm0.20}$ $\frac{1}{10.50\pm0.20}$ $\frac{1}{10.50\pm0.20}$ \frac	37 3.4 ± 0.6 0.6 ± 0.24 49.8 ± 0.37 43.6 ± 0.51 2.2 ± 0.37 $4.0^{\pm0.51}$ 51 1.8 ± 0.2 2.2 ± 0.37 46.6 ± 0.71 52.2 ± 0.37 2.6 ± 0.3 51 1.8 ± 0.2 2.2 ± 0.37 42.0 ± 0.71 52.2 ± 0.58 1.8 ± 0.2 3.6 ± 0.3 roups. roups. roups. roups. roups. roups. $\frac{1}{1}$ $\frac{2}{2}$ $\frac{Age}{120}$ $\frac{1}{18\pm0.2}$ $\frac{1}{2.8\pm0.2}$ $\frac{3.6\pm0.3}{3.6\pm0.2}$ $\frac{1}{1}$ $\frac{2}{2}$ $\frac{3}{3}$ $\frac{1}{12.55}$ $\frac{1}{2.2\pm0.58}$ $\frac{1}{1.8\pm0.2}$ $\frac{4.0^{\pm0.61}}{3.6\pm0.2}$ $\frac{1}{3.6\pm0.3}$ $\frac{1}{1}$ $\frac{2}{2}$ $\frac{3}{3}$ $\frac{1}{12.55}$ $\frac{1}{2.5\pm5.85}$ $\frac{1}{3}$ $\frac{1}{12.55}$ $\frac{1}{2.55}$ $\frac{1}{3}$ $\frac{1}{12.55}$ $\frac{1}{2.55}$ $\frac{1}{3}$ $\frac{1}{2.55}$ $\frac{1}{3}$ $\frac{1}{2.55}$ $\frac{1}{3}$ $\frac{1}{2.55}$ $\frac{1}{2.56}$ $\frac{1}{2.56}$ $\frac{1}{2.56}$ $\frac{1}{2.56}$ $\frac{1}{2.56}$ $\frac{1}{2.56}$ $\frac{1}{2.56}$ $\frac{1}{2.50\pm0.25}$ $\frac{1}{2$	37 3.4 ± 0.6 0.6 ± 0.24 $49.8^{\pm}-0.37$ 43.6 ± 0.24 2.8 ± 0.37 $4.0^{\pm}\pm0.27$ $4.0^{\pm}\pm0.37$ $4.0^{\pm}\pm0.31$ $4.0^{\pm}\pm0.31$ $4.0^{\pm}\pm0.31$ $4.0^{\pm}\pm0.21$	37 3.4 ± 0.6 0.6 ± 0.24 $49.8^{\pm}\pm0.37$ $4.6.6^{\pm}\pm0.37$ $4.6.6^{\pm}\pm0.37$ $4.0^{*\pm}-0.21$ 3.6 ± 0.37 $3.6,6\pm0.37$ $3.0,6\pm0.24$ $3.0,0+0.24$ $3.0,0$

9.85±0.20 d

9.70±0.21 d

9.60±0.20 c

8.10±0.18 b

+

+

4

Group No.	Tre	atment			Age /	week		
	I.S	Vacc.	0	1	2	3	4	5
1	-	-	5.30±0.13	6.10±0.15	6.30±0.12	6.42±0.20	6.20±0.21	5.90±0.20
2	+	-		6.21±0.14	6.20±0.13	6.40±0.18	6.18±0.25	6.00 ± 0.22
3	-	+			6.50±0.17	6.48±0.19	6.51±0.23	6.25±0.20
4	+	+			6.40 ± 0.16	6.41±0.20	6.48±0.25	6.20±1.54

Table (10): Effect of immunostimulant on uric acid in sera of vaccinated and non-vaccinated chickens with NDV live vaccine.

Table (11): Effect of immunostimulant on creatinine level in sera of vaccinated and non-vaccinated chickens with NDV live vaccine.

Group No.	Trea	atment			Age /	week		
	I.S	Vacc.	0	1	2	3	4	5
1	-	-	1.12±0.10	1.05 ± 0.03	1.15±0.03	1.25 ± 0.04	1.30±0.06	1.28±0.04
2	+	-		1.10 ± 0.04	1.10 ± 0.04	1.20 ± 0.04	1.26 ± 0.03	1.30 ± 0.04
3	-	+			1.21±0.05	1.25 ± 0.05	1.29 ± 0.05	1.40 ± 0.05
4	+	+			1.20 ± 0.03	1.23 ± 0.05	1.31±0.06	1.34±0.06

Table (12): Daily distribution of morbidity and mortality in challenged chickens.

Crown	Treatment		_				Ι	Days	post	-cha	lleng	ge			Total	%	
Group No.	I.S	Vacc.	Observation	1	2	3	4	5	6	7	8	9	10	11- 21	-		
1			Diseased No				5	7	4	2	1	-	-	-	19	95	
1	-	-	Died No.			3	4	5	5	2	1	-	-	-	20	100	
2	+		Diseased No				2	3	5	3	2	2	1	-	18	90	
2	+	-	Died No.				1	3	4	5	2	1	1	-	17	85	
3				Diseased No				1	1	2	1	1	-	-	-	6	30
3	-	+	Died No.					1	2	-	-	-	-	-	3	15	
4			Diseased No					1	1	1	1	-	-	-	4	20	
4	+	+ +	Died No.						1	-	-	-	-	-	1	5	

 Table (13): Results of VVND challenge test in immunostimulant medicated on vaccinated and non-vaccinated chickens.

Group No -	Trea	tment	- Total No of hirds	No of dood binds	No of survived birds	Drotaction 0/
Group No	I.S	Vacc.	- I otal ivo of birus	No of dead birds	No of survived birds	Frotection 70
1	-	-	20	20	0	0
2	+	-	20	17	3	15
3	-	+	20	3	17	85
4	+	+	20	1	19	95

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التأثير المناعى لمادة البيتا جلوكانز والمانان اوليجو سكاريدز على الكتاكيت المحصنة بلقاح النيوكاسل

تم دراسة التأثير المناعي لمادة البيتا جلوكانز والمانان اوليجو سكاريدز في الكتاكيت المحصنة بلقاح النيوكاسل وأظهرت النتائج أن الكتاكيت التي تم معاملتها بهذه المعاملات أعطت معدل أعلى في أوزان الجسم والغدة التيموسية والطحال وغدة فابريشيس كما أظهرت المجموعة المعاملة بهذه المواد مستوى أعلى في الكم والنوع في كرات الدم البيضاء وأظهرت القياسات للمجموعة المعاملة مستوى اقل من المجموعات الاخرى في إنزيمات الكبد كما أظهرت النتائج ارتفاع مستوى الأجسام والغدة التيموسية والطحال وغدة فابريشيس كما أظهرت معدل حماية العاملة بهذه المواد مستوى أعلى في الكم والنوع في كرات الدم البيضاء وأظهرت القياسات للمجموعة المعاملة معدومات الاخرى في إنزيمات الكبد كما أظهرت النتائج ارتفاع مستوى الأجسام المناعية للقاح النيوكاسل في هذه المجموعة مما أعطى معدل حماية اعلي في اختبارا لتحدى. من النتائج يمكننا أن نستنتج أن استخدام البيتا جلوكانز والمانان يؤدى إلى تحسن في الأوزان وكذا معدلات الصد ضد عدوى التحدي بفيروس النيوكاسل شديد الضراوة والمنتشر في مصر.