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EFFECT OF SEPARATE-SEX REARING ON BROILER PERFORMANCE, LITTER QUALITY AND IMMUNE RESPONSE.

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SUMMARY

One thousand and five hundreds feather sexed day-old chicks (Cobb 500) were divided into three equal groups; males, females and straightrun (as hatched). The 3 groups were reared identically. Body weight was significantly higher in male group compared with either female or straight-run groups. Also, the feed conversion ratio of the male group was significantly lower than the female group. The litter of the male group showed significantly lower moisture content as compared with the female group. No significant differences were observed in the maternal antibodies (for both ND and IBD viruses) between separately grown males and females. However, the non-specific immune response to SRBCs was significantly higher in females than males.

INTRODUCTION

Broilers may be raised sex-separate in order to

meet certain market demands and possible economic benefits. One of the major advantages of sex-separate rearing is product uniformity, since males are about 20% larger than females of the same age. Also, males will respond to higher plane of nutrition for a longer time than females.

The economic implications of raising separate sex flocks have been discussed by Leeson and Summers (1980) and Veerapen (1996).

In boilers, not only the final body weight but also the net carcass yield as well as the internal organs are affected by sex (Edwards et. al., 1973 and Susbilla et. al., 1994). Birds' activities e.g. feeding, drinking and resting are also affected by sex (Lei and van Beek, 1997 and Fayed et. al., 1996).

The genetic selection of poultry for growth rate and feed efficiency is thought to be accompanied by a reduction in immune response (Hayyari et. al., 1996). Moreover, in domestic fowl, stressful social environments result in less antibody activi-

ing sheep red blood cells (SRBCs) (Gross and ty against a variety of particulate antigens, includ-Siegel, 1973 and Siegel and Latimer, 1975).

The present study was carried out to evaluate the mance, litter quality criteria and immune response effect of separate-sex rearing on broiler perforto SRBCs (as anon-specific immune stimulant)

MATERIAL AND METHODS

I- Experimental birds and Management;

One thousand and five hundreds day old chicks (Cobb 500) were feather sexed according to Anon., 1988 and divided into three equal groups. second group to female chicks and the third group to both males and females. The birds were kept in The first group was assigned to male chicks, the an open sided house at Animal and Poultry Management Research Center, Faculty of Veterinary Medicine, Cairo University.

The birds were floor reared on a 10-cm thick layer of clean chopped wheat straw litter and received a commercial broiler ration, 23% crude protein and 3100 Kcal. ME until the 21st day of age then shifted to 22% crude protein and 3200 Kcal. ME ration until marketing age.

ed using eye drop method: Hitchner B1 at 7 days, The following vaccination programme was adopt-La Sota at 14 days and IBD BLEN at 15 days. While inactivated ND vaccine was injected at 17

days. From the beginning of the third week, bird were sprayed weekly using Hitchner B1 vaccine

II- Performance:

Average body weight was determined weeklyby weighing a sample of 50 birds from each group The feed consumption was recorded weekly and feed conversion ratio was estimated.

III- Examination of Litter:

were collected weekly in clean polyethylene bag Composite samples from surface and deep litter and examined for determination of: 1- pH: This was done electrochemically in a IIII dilution in distilled water. 2- Moisture content: The moisture percent was calculated after drying a 100 gm sample to a coffstant weight at 110 oC (Parsons and Baker, 1985)

propriate dilution of 1 gm sample using standard 3- Total colony count: By plating 1ml of the ap plate count agar.

IV-Immunological studies:

Sample of 25 birds from each group on days 1,1 and 14.2. and 14of age. The collected serum was used to many was us Blood samples were collected from a random measuring maternal antibody levels against castle disease (ND) virus using hacmagglutinalions inhibition. inhibition (HI) test (Beard, 1980) and Infection bursal 11. bursal disease (IBD) virus using agar gel processe (IBD) tation test (AGPT) (Cullen and Wyeth, 1975).

On the 15th day of age, 30 birds from each group were individually inoculated with 1 ml of 10% suspension of sheep red blood cells (SRBCs) as an antigenic stimulant (Gross, 1986). Blood samples were collected from the inoculated birds on days 3, 7, 14, and 21 post inoculation and the antibody response to SRBCs was measured using haemagglutination test (HA).

RESULTS AND DISCUSSION:

The effects of separate sex rearing on the average body weight and feed conversion ratio is shown in Table (1) and Fig. (1 and 2). It has been observed that separately grown male chickens had significantly higher average body weight from the beginning of the fourth week until the end of the sixth week of age (995, 1,230, 1,601 g) as compared to either separately grown females (949, 1,190, 1,550 g) or the straight-run group (721, 1,209, 1,530) respectively.

Also, the feed conversion ratio of the male group was significantly lower than that of the female group during the fifth and sixth week of age (1.9 and 2.1 compared to 2.1 and 2.3 respectively). These findings are in agreement with the other reports of Leeson and Summers (1980), Veerapen (1996) and Fayed et. al., (1996).

Concerning litter quality criteria, the obtained results (Table 2 and Fig. 3) revealed no significant differences in either litter pH or total colony counts between separately grown sexes and straight-run group. However, the litter pH showed a characteristic shift to the alkaline side by the end of the 3rd week of age in the 3 experimental groups. Also, the total colony counts exhibited a grad and steady rise as the birds advanced in age.

As for the litter moisture %, it is noticed that, from the beginning of the third week of age, the female group showed a significantly higher moisture content (16.65, 12.65, 23.9 and 33.45%) as compared with either male group (12.5, 10, 17.15 and 28.4%) or straight-run group (8.3, 9.8, 14, 28) respectively. Dry litter in the male group could be possibly attributed to the differences in physical activities between males, females and straight-run birds (Fayed et. al., 1996). Males, being more active, provided better chances for maintaining dry litter.

Results in tables (3 and 4) show that screening of the maternal antibodies for both ND and IBD viruses on 1st, 7th and 14th days of age, revealed no significant differences between separately grown males and females. On the other hand, the non-specific immune response to SRBCs (Table 5 and Fig. 4) revealed significantly higher HA antibod-

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Table (1): Effect of separate-sex rearing on broiler performance.

Age (Weeks)	Males		Females		Straight run	
	Body Weight/g	FCR*	Body Weight/g	FCR	Body Weight/g	FCR
1	115.2	1.12	106	1.2	101	1.16
2	246	1.35	237	1.45	259	1.10
3	458	1.6	435	1.7	469	1.65
4	995	1.8	949	1.9	721	1.85
5	1.230*	1.9**	1.190	2.1	1.209	2
6	1.601*	2.1**	1.550	2.3	1.530	2.2

* Feed Conversion Ratio.

Significant differences P(0.05 in body weight between male and both female and straight-run groups

** Significant differences P(0.05 in FCR between male and female groups.

Table (2): Effect of separate sex rearing on litter quality.

Age (Weeks)	Males			Females			Straight run		
	Moisture %	рН	T.C.C*	Moisture %	рН	T.C.C*	Moisture %	рН	T.C.C
1 2	10 7.85	6.6 9	4.7 7.49	10 8.05	6.6	4.7	11 .	6.6	4.7
3	12.5	8.4	7.65	16.65*	8.6 8.4	6.75 7.88	6.6 8.3	9.6 8.5	6.9 6.9
5	10 17.15	7.9 7	8.79 8.4	12.65* 23.9*	8.3 7	8.18 8.72	9.8 14	8.4 7	8.15 8.9
6	28.4	7	9.65	33.45*	7	9.39	28	7	7.96

*Log total colony count/ g dry weight.

* Significant differences P≤ 0.05 in moisture % between female and both male and straight number of the straight number

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Table (3): Maternal antibodies against Newcastle disease in sera of chickens grown separately.

	Arithmetic mean of HI* test			
Time of testing	Males	Females		
Day old	3.6	3.6		
7 days	2.9	2.9		
14 days	3.2	3.2		

^{*} Haemagglutination inhibition.

Table (4): Maternal antibodies against infectious bursal disease in sera of chickens grown separately.

Time of the still	% of positives AGPT*		
Time of testing	Males	Females	
Day old	90%	90%	
7 days	60%	60%	
14 days	40%	60%	

^{*} Agar gel precipitation test.

Table (5): Haemagglutinin antibody response (GMT)° of separately reared chickens inoculated with sheep red blood cells (SRBCs) at 15 days of age.

Males	Females	Control
0		
	0	0
	2.5*	0
3.2	2.95	0
3.5	The second second	0
	0 1.9 3.2 3.5	0 0 1.9 2.5* 3.2 2.95

significant differences P(0.05 between female and male

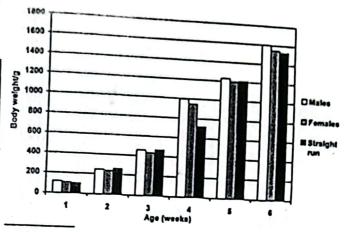


Fig.(1): Effect of separate sex rearing on body weight

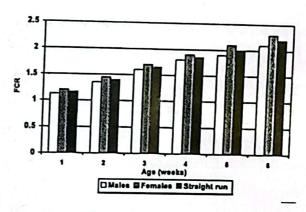


Fig.(2): Effect of separate sex rearing on feed conversion ratio (FCR)

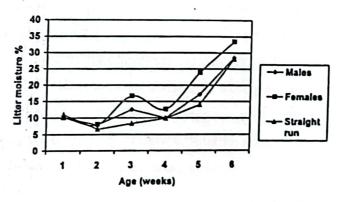


Fig.(3): Effect of separate sex rearing on litter moisture

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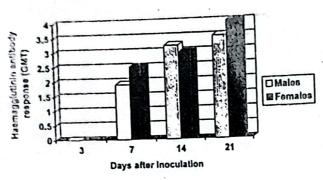


Fig.(4): Haemagglutinin antibody response of separately reared chickens inoculated with SRBCs at 15 days

ies in separately grown females on the 7th and 21st days post inoculation (2.5 and 4) as compared to separately grown males (1.9 and 3.5) respectively. The observed reduction in HA antibodies to SRBCs in separately grown males is certainly attributed to higher growth rate in males (Hayyari et. al., 1996).

Conclusively, the economic implications of raising separate sex flocks should be considered in relation to the cost of sexing day old chicks, and changes in processing or marketing strategies. According to the present study, the difference in immune response to SRBCs between males and females may suggest a different schedule of vaccination against common poultry pathogens. However, this point needs further investigation.

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