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Performance, carcass quality, blood parameters and Immune System of broilers fed diets supplemented with oregano oil (*Origanum* sp.)

Navid Hosseini Mansoub

Department Of Animal Science, İslamic Azad University, Maragheh Branch, Iran

ABSTRACT

Present experiment was performed to study the effects of oregano oil (Origanum sp.)medicinal plant on performance, blood biochemical and immunity parameters of broiler chickens. During the experiment 400 chicken broilers were divided in four experimental groups with five repetitions: control group (C) without any oregano oil, group 1 (G1) received 100 ppm of oregano oil, group 2 (G2)with 150ppm of oregano oil, grop3(G3) received 200 ppm of oregano oil. From 1 day to 42 days, the highest amount of body weight gain and the lowest level of FCR were observed in the groupG3 but the best result for daily feed intake was in G2 and the lowest group was observed in control group. The lowest percent of abdominal fat was observed in experimental group 2 and the highest percent of breast was in experimental G3. The results showed that using oregano oil (Origanum sp.) in chickens diet had not significant effects on on blood biochemical parameters and immune system of broiler chickens (p>0.05).

Keywords: Oregano oil, Broilers, Carcass traits, Blood biochemical.

INTRODUCTION

There are a lot of reports indicating the positive effects of herbs like anti-coccidal, anti-oxidant, anti-fungi and etc. Some of medical effects of herbs are related to their secondary metabolites such as phenols, necessary oils, saponins and etc [1].Consequently there is considerable research interest in the possible use of natural products, such as essential oils and extracts of edible and medicinal plants, herbs and spices, for the development of new additives in animal feeding. Aromatic plants like Sea-buckthorn contain flavonoids described previously as stimulators of the immune response, these dietary flavones having an effect against microbial infection [2-3].The antimicrobial activity of essential oils derived fromspices and herbs [4-5] is of interest as these oils could be used asfeed additives alternative to antibiotics [6].

Lots of studies on phytogenic compounds of plants essential oils have been performed while there are limited evidences about the effect of herbal solid forms on live birds health and performance. Easy and practical application, availability and less cost are known as advantages of the whole herbs application in compare to extracted or essential oil forms. In the other hand, a synergistic effect of phytogenic compounds have been reported in studieswith essential oils [7]. A recent study involving live birds showed thatblends of the primary components of the essential oilscould be used to control *Clostridium perfringens*, thebacterium that causes necrotic enteritis in broilers. Ground thyme has been shown toinhibit the growth of *S. typhimurium*when added tomedia [8]. The main functions of the essential oils cover pathogen control including antimicrobial activity [9], antioxidant activity [10], digestion aid including stimulation of endogenous enzyme activity and nitrogen absorption [11] and inhibition of odour and ammonia control [12].

Therefore, present experiment was planned to study the effects of oregano oil (Origanum sp.)on performance, carcass quality, blood biochemical parameters and immunity parameters of broilers chickens.

MATERIALS AND METHODS

During the experiment 400 chicken broilers were divided in four experimental groups with five repetitions: control group (C) without any oregano oil, group 1 (G1) received 100 ppm of oregano oil, group 2 (G2)with 150ppm of oregano oil, grop3(G3) received 200 ppm of oregano oil. unbound water and dietary was in poultries' access. Dietary and chick weigh were going on weekly. Feed consumed was recorded daily, the uneaten discarded, and feed conversion ratio (FCR) was calculated (total feed : total gain). At the end of experiment, some analyses was done via SAS (Statistical Analyses Software) in the statistical level of 5% according to data gathered from dietary, weight improvement, average of FCR, weight of rearing period and carcass yield. At 42 days of age, four birds per replicate were randomly chosen, slaughtered and carcass percent to live weight and percent of carcass parts to carcass weight were calculated.

In the 35th day of experiment, three chicks were chosen from each group and inoculated from brachial vien by 0.1 ml (5 %). Heterophils to Lymphocytes ratio were determined which had been obtained from barchial vein of three randomly chosen chicks from each group in the 42th day of experiment.Blood samples were obtained from barchial vein and centrifuged in order to getting serum, after 12 hours of fasting in the 42th day of experiment.

Ingredients	Starter	Grower
Maize	559	296
Wheat		334
Soybean meal	368	298
Soybean oil	32	42
Fish meal	18	
Limestone	10	
Oyster shell		12
Dicalcium phosphate	5	15
Vitamin-mineral mix	5	5
dl-methionine	1	1
Sodium chloride	2	2
Vitamin E (mg/kg)		98
Zn		52
Analyzed chemical composition (g/kg)		
Dry matter	891.9	893.1
Crude protein	221.1	200.8

 Table 1. Ingredients and chemical analyses composition of the starter and grower diets

Navid Hosseini Mansoub

Fat	62.3	63.1
Fiber	36.1	35.6
Ash	61.7	57.0
Calcium	8.22	8.15
Phosphorus	5.45	5.39
Selenium (mg/kg)	0.58	0.56
ME by calculation (MJ/kg)	12.69	12.71

vitamin A, 9,000 IU; vitamin D3, 2,000, IU; vitamin E, 18 IU; vitamin B1, 1.8 mg; vitamin B2, 6.6 mg B2,; vitamin B3, 10 mg; vitamin B5, 30 mg; vitamin B6, 3.0 mg; vitamin B9, 1 mg; vitamin B12, 1.5 mg; vitamin K3, 2 mg; vitamin H2, 0.01 mg; folic acid, 0.21 mg; nicotinic acid, 0.65 mg; biotin, 0.14 mg; choline chloride, 500 mg; Fe, 50 mg; Mn, 100 mg; Cu, 10 mg; Zn, 85 mg; I, 1 mg; Se, 0.2 mg.

RESULTS AND DISCUSSION

Table 2 shows the effect of different dietaryoregano oil (Origanum sp.)on performance of boiler chickens. According to comparisons of this table it has been proven that highest amount of body weight gain and the lowest level of FCR were observed in the group G3 but the best result for daily feed intake was in G2 (p<0.05).

The beneficial effect of growth promoting feed additives on animals arisesfrom stabilizing feed hygiene and beneficially modulating the gut ecosystemby controlling potential pathogens. Phytogenic compounds have a number ofactive ingredients and pharmacologically active substances that are beneficial for maintaining health and improving performance of poultry and other livestockspecies. They are reported to stimulate secretion of digestive enzymes (lipase andamylase) and intestinal mucous in broilers, to stimulate feed digestion, to impairadhesion of pathogens and to stabilize microbial balance in the gut [13].

Table 3 shows the effect of plants and their different combinations on carcass and it's parameters. According to the data, there are significant differences in the carcass characters (p<0.05). The lowest percent of abdominal fat was observed in experimental group 2 and the highest percent of breast was in experimental G3. Aromatic plants and essential oil extracted from these plants have been used as alternatives to antibiotics. For this reason, these plants are becomingmore important due to their antimicrobial effects and the stimulating effect on animal digestive system[14]. The active principles of essential oils act as a digestibility enhancer, balancing the gut microbial ecosystem and stimulating the secretion of endogenous digestive enzymes and thus improving growth performance in poultry [15]. The effects of experimental plants on blood biochemical parameters are presented in Table 4. there is no effect on blood biochemical parameters and immune system of broiler chickens

Experiment	Weigh	Food	Average	Average
Treatments ¹	Improvement	Intake (G)	of FCR	Of Weight
С	40.9 ^a	84.9 ^a	1.81 ^a	1998.9 ^a
G1	41.1 ^a	85.3 ^a	1.80^{a}	1999.4 ^a
G2	41.3 ^a	86.6^{ab}	1.70^{ab}	2000.7^{a}
G3	41.8^{ab}	86.1 ^{ab}	1.61^{ab}	2005.3 ^{ab}
SEM	0.98	1.12	0.02	28.6
P-value	0.02	0.006	0.003	0.02

 Table 2: Effects of treatments on performance of broilers.

a-c Means with in columns with different superscript differ significantly

Parameters	C1	G1	G2	G3	SEM
Abdominal Fat	3.86 ^a	3.74 ^a	3.53 ^{ab}	3.60^{ab}	0.35
Gizzard	3.08	3.10	3.15	3.25	0.11
Breast	33.18 ^b	33.09 ^b	34.20^{b}	35.24^{ab}	1.52
Thigh	26.11	25.48	25.43	28.23 ^a	1.02
Liver	3.27 ^a	2.90 ^a	3.62 ^a	4.14^{ab}	0.33

Table 3. The effect of different levels of Oregano oil on carcass traits of broilers

Means with different subscripts in the same column differ significantly (P < 0.05)

Table 4: Effect of different levels oregano oil on immune system parameters of broiler chickens.

Parameters	C1	G1	G2	G3	SEM
Heterophils to Lymphocytes ratio	0.20	0.22	0.20	0.21	0.03
Globulin	1.40	1.41	1.44	1.43	0.19
Albumin	1.40	1.53	1.44	1.55	0.12

 $^{a-c}$ Means with different subscripts in the same row differ significantly (P < 0.05)

Table 5. The effect of different levels oregano oil on blood biochemical of hens

Blood Parameter	С	G1	Treatments G2	G3	SEM
Glucose (mg/dl)	170.36	171.03	171.42	173.30	1.46
Cholesterol (mg/dl)	134.60	135.02	135.22	134.90	2.12
Triglyceride (mg/dl)	42.60	41.97 ^a	41.60	42.10	1.80
LDL	33.13	33.29	32.19	32.69	1.02
HDL	82.22	82.65	83.16	83.29	1.55

Means with different subscripts in the same column differ significantly (P < 0.05)

REFERENCES

[1] M.A. Ipu, M.S. Akhtar, M.I. Anjumi, and M.L. Raja., *Pakistan Veterinary Journal*.2006, 26,144-148.

[2] KAMEL C. Tracing modes of action and the roles of plant extracts in non-ruminants. Pages 135-150 in Recent Advances in Animal Nutrition.P.C.Garnsworthyand J. Weiseman, ed. Nottingham University Press, Nothingham, UK. **2001**.

[3] Mishra KP, Chanda S, Karan D, Ganju L, Sawhney RC. Phytother Res. 20, (2008).

[4] Deans, S.G. and G. Ritchie, 1987. Int. J. Food Microbiol., 5: 165-180.

[5] Hammer, K.A., C.F. Carson and T.V. Riley, **1999**. J. Appl. Microbiol., 86: 985-990.

[6] Taylor, D.J., 2001. Br. Poult. Sci., 42: (Suppl) 67-68.

[7] Mitsch P, Zitter-Eglseer K, Kohler B, Gabler C, Losa R, Zimpernik I (2004). *Poult. Sci.*, 83: 669-675.

[8] Aktug, S. E. and M. Karapinar, 1986. Intern. J. Food Microbiol., 3: 349-354.

[9] Cowan, M.M., 1999. Clin. Microbiol. Rev. 12, 564-582.

[10] Botsoglou, N.A., Christaki, E., Florou-Paneri, P., Giannenas, I., Papageorgiou, G. & Spais, A.B., **2004**. *Afr. J. Anim. Sci.* 34, 52-61.

[11] Gill, C., 2001. Feed Int. 22 (3), 40-45.

[12] Varel, V. H., 2002. J. Anim. Sci. 80 (2), E1-E7.

[13] Lee, K.W., Everts, H. and Beyen, A.C. **2003**. *Journal of Applied Poultry Research*. 12:394-399.

[14]Osman, N., G. Talat, C. Mehmet, D. Bestami and G. Simsek, 2005. Intern. J. Poult. Sci., 4: 879-884.

[15] Lovkova M.Y., Buzuk G.N., Sokolova S.M., Kliment'eva N.I. 2001: Appl. Biochem. Microbiol., 37, 229–237.