

Influence of Different Levels of Phytase Enzyme on Japanese Quail (*Coturnix japonica*) Performance

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Summary

An experiment was conducted to study the effect of phytase enzyme (Natuphos 500) supplementation in Japanese quail (*Coturnix japonica*) (3 to 45 days of age) on body weight, carcass performance, percentage of tibia bone ash and tibia phosphorus. Data were analyzed as a completely randomized design arranged with five levels (150, 300, 600, 1200 and 2400 FTU/Kg) of phytase enzyme as treatments 1, 2, 3, 4 and 5, respectively, and two control groups (positive and negative controls). None of the phytase levels had effect on carcass yield at 45 days of age ($p>0.05$), except first treatment with 70.06% of carcass yield that was significantly different than treatment 5 ($p<0.01$). The treatment 5 for percentage of tibia ash was 59.83% that was significantly different compared to the other treatments ($p<0.01$). Treatment 3 increased ($p<0.01$) phosphorus of tibia ash at 45 days of age (240.22 mg). Experimental replicates were not significantly different in any stage ($p>0.05$). Female quails at body weight, slaughter weight, carcass weight without viscera and legs weight were better than male ($p<0.01$), but male quails at carcass yield (69.06%) were significantly different ($p<0.01$) in contrast to the females (66.30%).

Key words

Japanese quail, phosphorus, phytase enzyme

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Introduction

Approximately two-thirds of the total phosphorus (P) in plants, which are the major constituents of poultry diets, is in the form of phytate (Punna and Roland, 1999; Viveros et al., 2000) and it is unavailable or poorly available when utilized by humans and other monogastric animals. This unavailability is due to the very low phytase activity found in the digestive tract (Pallauf et al., 1994). Therefore, diets of monogastric animals are often supplemented with sources of inorganic P to meet the P requirements of the animal, which increases the cost of the diets and contributes to environmental pollution. Dietary supplementation with sources of microbial phytase is well established as an effective and practical method of improving phytate digestibility in production animals. Phytase is an enzyme that hydrolyzes phytate to inositol and inorganic phosphate. Phytases are naturally found in a number of seeds including cereals, legume, by-products, and other feedstuffs (Eeckhout and De Paepe, 1994; Viveros et al., 2000) and in microbial sources (Wyss et al., 1999).

Nelson et al. (1968) demonstrated that the addition of the enzyme phytase to grains and feeds was an effective way to increase phosphorus availability to poultry. Numerous reports have shown that phytate phosphorus present in cereal grains could be made available to both poultry and swine by treating the grains or supplementing the diets with the enzyme phytase, reducing the need to supplement these diets with inorganic phosphorus and decreasing phosphorus excretion (Nelson et al., 1971; Jongbloed and Kemme, 1990; Simons et al., 1990; Jongbloed et al., 1992; Ketaren et al., 1993; Mitchell and Edwards, 1996a, b). The commercial development of phytase has provided the animal industry with a tool to decrease the need to supplement animal feeds

with inorganic phosphorus, which will decrease phosphorus in animal manures, and consequently decrease phosphorus loading of agricultural lands.

Phytase research that has focused on the effect of varying levels of supplemental phytase has shown increases in broiler performance, bone ash, and phytate P utilization with each additional unit or level of phytase (Shirley and Edwards, 2003).

Most of the investigations on phytase are with chickens while quails were less considered. Therefore, the target of this study is the investigation of different levels of phytase enzyme (Natuphos) on Japanese quail (*Coturnix japonica*) performance.

Materials and methods

A total of 168, 3-d-old Japanese quail chicks, (*Coturnix japonica*) were housed in pen with the newspaper litter until 12 days of age. Then birds were moved into one-tier cage (60×50×40 cm) in an environmentally controlled room, with 23-h constant overhead fluorescent lighting. The temperature was decreased gradually from 35°C to 20°C on the weekly bases. The chicks were allocated to 21 cages, each cage containing 8 chicks that were receiving seven diets with three replication of each diet. The experimental diets are presented in Table 1 and were formulated to meet the requirements of quail as established by the NRC (1994). Experimental diets were arranged with five levels 150, 300, 600, 1200 and 2400 phytase enzyme unit (FTU/Kg) with commercial name (Natuphos 500) as treatments 1, 2, 3, 4 and 5, respectively, and two control groups (positive and negative controls). One FTU is defined as the amount of enzyme that liberates micromole inorganic phosphorus per minute from 0.0051 mol/L sodium phytate at 37°C and at pH = 5.5 (Hall et al.,

Table 1. Ingredients and nutrient content of the experimental diets

Diets Ingredients (%)	0 (positive control)	0 (negative control)	Phytase Levels (FTU/Kg) ¹				
			150	300	600	1200	2400
Corn (8.5% CP)	49	49	49	49	49	49	49
Soybean meal(44% CP)	44.91	44.91	44.91	44.91	44.91	44.91	44.91
Vegetable oil (8.8 kcal/g)	2.865	2.865	2.865	2.865	2.865	2.865	2.865
Limestone	1.304	1.304	1.304	1.304	1.304	1.304	1.304
Salt (NaCl)	0.20	0.20	0.20	0.20	0.20	0.20	0.20
Dicalcium phosphate	0.747	0.212	0.212	0.212	0.212	0.212	0.212
DL-Methionine	0.136	0.136	0.136	0.136	0.136	0.136	0.136
Mix vitamins ²	0.25	0.25	0.25	0.25	0.25	0.25	0.25
Trace mineral ³	0.25	0.25	0.25	0.25	0.25	0.25	0.25
Sand	0.337	0.873	0.87	0.867	0.861	0.849	0.825
Phytase (5000 FTU/ g)	0	0	0.003	0.006	0.012	0.024	0.048
Total	100	100	100	100	100	100	100
Nutrient (Calculated)							
Metabolizable energy (Kcal/Kg)	2900	2900	2900	2900	2900	2900	2900
Crude protein (%)	24	24	24	24	24	24	24
Methionine + cystine (%)	0.884	0.884	0.884	0.884	0.884	0.884	0.884
Lysine (%)	1.33	1.33	1.33	1.33	1.33	1.33	1.33
Calcium (%)	0.800	0.682	0.682	0.682	0.682	0.682	0.682
Total phosphorus (%)	0.569	0.469	0.469	0.469	0.469	0.469	0.469
Available phosphorus (%)	0.300	0.200	0.200	0.200	0.200	0.200	0.200

¹ Danisco Animal Nutrition, Carol Stream, IL, FTU/kg feed = phytase units/kg feed. One phytase unit (FTU) is defined as the amount of enzyme that liberates micromole inorganic phosphorus per minute from 0.0051 mol/L sodium phytate at 37°C and at pH = 5.5; ² Provided per kilogram of diet: vitamin A (as retinyl acetate), 8,000 IU; cholecalciferol, 2,200 ICU; vitamin E (as DL- α -tocopheryl acetate), 8 IU; vitamin B12, 2800.02 mg; riboflavin, 5.5 mg; D-calcium pantothenic acid, 13 mg; niacin, 36 mg; choline, 500 mg; folic acid, 0.5 mg; vitamin B1 (thiamin mononitrate), 1 mg; pyridoxine, 2.2 mg; d-biotin, 0.05 mg; vitamin K (menadione sodium bisulfate complex), 2 mg; ³ Trace mineral mix provided the following (per kg of diet): manganese (MnSO₄·H₂O), 60 mg; iron (FeSO₄·7H₂O), 30 mg; zinc (ZnO), 50 mg; copper (CuSO₄·5H₂O), 5 mg; iodine (ethylene diamine dihydroiodide), 1.5 mg.

2003). Small amounts of the basal diet were first mixed with the respective amounts of phytase as a small batch and then with a larger amount of the basal diets until the total amounts of the respective diets were homogeneously mixed. Diets in mash form and water were provided ad libitum from 3 to 45 days of age.

At the age of 45 days, four chicks were randomly selected from each treatment and weighed, and were killed for carcass performance in agreement with the method of Yalcin et al. (1999). Three quails were randomly selected from each treatment and a total of 21 left legs were banded for identification percentage tibia ash (TA) and phosphorus samples of bones tibia in agreement with the method of AOAC (1990). Data were subjected to analysis of variance using the general linear models procedures of SAS software (SAS Institute, 1990). Statements of statistical significance are based on a probability of $P \leq 0.05$. The statistical model used was:

$$Y_{ijk} = \mu + T_i + S_j + TS_{ij} + R_k + e_{ijk}$$

where Y_{ijk} is the individual observation, μ is the overall mean, T_i is the treatment effects ($i=1, 2, \dots, 7$), S_j is the sex effects ($j=1, 2$), TS_{ij} Interaction effect between i^{th} dietary treatment and j^{th} sex, R_k is the replication effects, and e_{ijk} is the error term.

Result and discussion

The effects of different levels of dietary phytase on the body weight (BW), slaughter weight (SW), carcass weight without viscera (CWV), carcass yield (CY), percentage of tibia bone ash (Tash) and tibia phosphorus (TP) of quails at 45 days of age are summarized in Table 2. The data indicated that body weights at 45 days of age were affected by phytase enzyme ($p < 0.01$). So that quails of treatment 5 significantly increased body weight (195.16 g) compared to those on the negative control group that had weight 173.90 g. These results were nearly to those obtained by Huff et al. (1998), Viveros et al. (2002) and Hall et al. (2003) in broiler. Likewise sex was a significant source of variation for body weights at 45 days of age ($p < 0.01$). Females showed 196.12 and male quails showed 169.74 g. The superiority of female versus male agreed with previous studies of Vali et al. (2005) and Vali

et al. (2006). It is necessary to notify that the hydrolysis and absorption of phytate phosphorus (P) by monogastric animals are complex processes that are influenced by many factors. Dietary ingredients and feed processing seem to be the most important factors related to the diet, while age and type of birds could also affect phytate (P) utilization (Reddy et al., 1982; Sebastian et al., 1998; Attia et al., 2003). This possesses a problem to nonruminant animals because they do not produce sufficient amounts of intrinsic phytase which is necessary to hydrolyze the phytic acid complex EI-Badry et al. (2008).

The effects of experimental replications 1, 2 and 3 on body weight at 45 days of age 180.04, 184.90 and 181.73 g, respectively were not significantly different ($p > 0.05$).

Carcass yield of quails treatment 1 was 70.06% and treatment 5 was 66.73% that was significantly different ($p < 0.01$), but treatment 1 had not significant difference ($p > 0.05$) with the rest of treatments. Also carcass yield of male quails with 69.06% had better performance than females (66.30%) that was significantly different ($p < 0.01$).

These results demonstrated that treatment 3 had significantly different percentage of tibia ash (59.83%) compared with the rest of treatments ($p < 0.01$), but in comparison with positive control (57.72%) there was no significant difference ($p > 0.05$). This increment has been reported by several authors working with chickens and it is considered a good indication of bone mineralization (Broz et al., 1994; Qian et al., 1996; Sebastian et al., 1996a, b). The improvement in ash percentage in tibia probability can be related to the increase in Ca, P, Mg, and Zn retentions from the phytate-mineral complex by the action of phytase. In contrast, Keshavarz (2000) observed in pullets and Rama-Rao et al. (1999) in chickens that tibia ash was not influenced by phytase in the diets.

These results demonstrated that quails treatments 2 and 3 improved performance of P in tibia 217.38 and 240.22 mg respectively, and they were significantly different compared to other treatments ($p < 0.01$).

Table 2. Influence of dietary phytase levels (FTU) on the body weight at 45 days of ages (BW), slaughter weight (SW), carcass weight without viscera (CWV), carcass yield (CY), percentage of tibia bone ash (Tash) and tibia phosphorus (TP) of Japanese quails (*Coturnix japonica*)

Main effects		BW (g)	SW (g)	CWV (g)	Cy (%)	Lw (g)	Tash (%)	TP(mg)
Overall mean		182.18	173.31	123.13	67.75	32.23	55.75	171.34
Treatment	1	175.25 ^{abc}	166.39 ^{bc}	122.71 ^{ab}	70.06 ^a	30.58 ^{bc}	53.61 ^c	132.46 ^d
	2	177.71 ^{abc}	169.38 ^{abc}	122.12 ^{ab}	68.83 ^{ab}	32.65 ^{abc}	55.54 ^{bc}	217.38 ^{ab}
	3	190.00 ^{abc}	178.48 ^{abc}	127.28 ^{ab}	67.27 ^{ab}	33.36 ^{ab}	59.83 ^a	240.22 ^a
	4	170.62 ^c	161.62 ^c	115.45 ^b	67.72 ^{ab}	29.13 ^c	55.60 ^{bc}	164.94 ^{cd}
	5	195.16 ^a	186.91 ^a	130.00 ^a	66.73 ^b	34.74 ^a	55.62 ^{bc}	172.07 ^c
Control ¹	+	191.12 ^{ab}	182.62 ^{ab}	126.14 ^{ab}	66.14 ^{ab}	33.09 ^{abc}	57.72 ^{ab}	190.45 ^{bc}
	-	173.90 ^{bc}	165.79 ^{bc}	116.53 ^{ab}	67.12 ^{ab}	31.80 ^{abc}	52.36 ^c	81.93 ^e
Replication	1	180.04 ^a	171.35 ^a	121.76 ^a	67.68 ^a	31.57 ^a	55.10 ^a	185.24 ^a
	2	184.9 ^a	175.81 ^a	124.71 ^a	67.71 ^a	32.63 ^a	55.61 ^a	163.63 ^a
	3	181.73 ^a	172.88 ^a	123.04 ^a	67.87 ^a	32.6 ^a	56.54 ^a	165.17 ^a
Sex	f	196.12 ^a	186.10 ^a	129.65 ^a	66.30 ^b	34.33 ^a	55.88 ^a	175.55 ^a
	m	169.74 ^b	161.64 ^b	117.19 ^b	69.06 ^a	30.32 ^b	55.58 ^a	165.74 ^a

¹ Positive and negative control groups; ^{abcde} Values with in a column with no common superscript differ significantly ($p \leq 0.05$).

Data also showed that carcass yield, tibia ash, and tibia phosphorus were not affected at any stage by experimental replications ($p>0.05$; Table 2).

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