Influence of Dietary Phytase, Lysine and Phosphorus on Nutrient Digestibility, Hydrolysis of Phytic Acid and Coenzyme Q10 Synthesis in Laying Hens

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Abstract

High doses of phytase have been shown to increase phytic acid (Phyt-ac) hydrolysis, nutrient availability and coenzyme Q10 (CoQ10) synthesis in broilers, but no previous reports are available in layers. The present study tested the effects of four doses of phytase (0, 300, 1200 and 4800 FTU/kg) and three digestible lysine (dLys) levels (0.67, 0.77, and 0.87%) on Phyt-ac hydrolysis, amino acid and mineral digestibility in layers fed diets with 0.12% available P (avP). Additionally, a diet with 0.25% avP and 0.87% dLys, without phytase added, was evaluated as control treatment. CoQ10 content in liver was quantified in the first three doses (0, 300, and 1200 FTU/kg) at the highest and lowest levels of dLys (0.67 and 0.87%). There was an interaction effect (P<0.05) between phytase doses and dLys levels for amino acid digestibility. In 0.67% dLys diets, the response to incremental doses of phytase was linear, while in 0.77% and 0.87% dLys there was a quadratic response (P<0.05). P digestibility was higher with incremental doses of phytase, despite a reduced effect of enzyme in 0.87% dLys diets (P=0.034). No differences (P>0.05) were observed between treatments on availability of Ca, Zn, Cu, Co, Mn and Fe. Phytase inclusion increased Mg, Na, and K digestibility in 0.67 and 0.77% dLys diets, but in 0.87% diets these were reduced (P<0.05). Doses of phytase 1200 and 4800 FTU/kg increased the presence of lower low inositol esters (Ins-P) such as Ins-5P, Ins-3P and inositol (P<0.05). Ins-4P and Ins-2P content was influenced (P<0.05) by phytase addition and dietary dLys. Liver CoQ10 content was similar between phytase doses in 0.67% dLys diets, but in the 0.87% dLys diet, phytase addition reduced coenzyme concentration in liver (P<0.05). In conclusion, phytase dose and dLys level in layer diet affected nutrient digestibility, Phyt-ac hydrolyzes, and CoQ10.

Keywords: Phytase superdosing; Dietary digestible lysine; Ileal digestibility; Hydrolysis of phytic acid; Coenzyme Q10 in liver; Laying hens

Introduction

The capability of phytic acid (Phyt-ac) to impact the availability of nutrients is due to the presence of 12 replaceable protons with the capacity to associate with other molecules [1]. These chelates in the gastro-intestinal lumen can reduce the absorption and utilization of nutrients such as proteins, amino acids and minerals [2]. Otherwise, the anti-nutritive effect of Phyt-ac can be eliminated by hydrolysis of Phyt-ac covalent bonds by phosphatases, including phytases. During the hydrolysis process are obtained low inositol phosphates (Ins-P), mono-phosphate and free myo-inositol [3]. Previous studies have shown that lower Ins-P esters can negatively affect protein digestibility by limiting solubility and/or blocking digestive enzymes in the gastro-intestinal tract [4].

The extent of phytate hydrolysis in the gut will depend on the dietary phytase level. Walk CL, et al. [5] found that using 1500 FTU/ kg, Ins-5P, Ins-4P and Ins-3P content is reduced 33% compared with 500 FTU/kg, and this last dose is less effective in realizing nutrients. For that, superdosing could highly impact the availability of nutrients, primarily P, but also for protein and mineral [6].

The extra-phosphoric effect of phytase implicates changes in metabolic pathways, modifying metabolic system including coenzymes [7]. Coenzyme Q10 (CoQ10) is recognized as crucial component in the energy producing processes, and an endogenous antioxidant in the cell as well. In the inner mitochondrial membrane, CoQ10 is an electron carrier in the respiratory chain for the formation of ATP. In the complex III of the respiratory chain, CoQ10 is present in three forms: ubiquinone (oxidized-coQ10), ubisemiquinone, and ubiquinol (reduced-CoQ10H$_2$) [8]. Karadas F, et al. [7] found higher CoQ10 levels in the liver using 12500 FTU/kg of a 6-phytase derived from *Escherichia coli* or with 0.45% avP compare to 0.25% avP in 21-day old broilers fed with or without 250 FTU/kg phytase.

Another variable that can influence the degradation of Phyt-ac is the P content. Hughes A, et al. [9] reported a negative linear response in phytate ileum digestibility with 0.25% avP content in diet when testing 200, 400, and 600 FTU/kg phytase; however, reducing avP to 0.15%, no linear response was observed. In the same study, 500 FTU/kg increased phytate digestibility in 0.15% avP compare to 0.35% avP. Agbede JH, et al. [10] found higher content of Ins-6P with a dietary content of 0.8% total P than 0.4% in broilers.

Lysine requirements for dietary content have been evaluated in laying hen [11], although there is limited ileal digestibility data available, with no previous evaluation of the impact of phytase superdosing with graded levels of digestible Lys (dLys).

In the present study was hypothesized that phytase superdosing (1200 and 4800 FTU/kg) would increase Phyt-ac hydrolysis, improving nutrient digestibility in ileum of laying hens fed a corn-soybean meal diet. Furthermore, was expected that coenzyme Q10 content in liver would be influenced by phytase dose, but not by dietary avP content. Thus, the present study evaluated the effect of inclusion of four doses of a exogenous phytase (0, 300, 1200, and 4800 FTU/kg) in 0.12% avP diets with three levels of dLys (0.67, 0.77, and 0.87%) on ileal digestibility of essential (except tryptophan) and non-essential amino acids, minerals (Zn, Fe, Mg, Cu, Mn, Ca, Na and K), and on Phyt-ac hydrolysis. Additionally, one diet with 0.25% avP and 0.87% dLys was included, with the aim to evaluate the impact of avP level on each one of the above-mentioned variables. The CoQ10 content in liver was quantified for three phytase doses (0, 300, and 1200 FTU/kg) and two dLys levels (0.67% and 0.87%).

Materials and Methods

Experimental design

Four doses of an exogenous phytase (0, 300, 1200 and 4800 FTU/kg) were tested with three levels of dLys (0.67, 0.77 and 0.87%) in a corn-soybean meal based on diet containing 0.12% avP. The control phytase-free diet with 0.25% of avP and 0.87% dLys level was evaluated as well. With the exception of dLys and avP, diets were formulated following the NRC (1994) recommendations for white-line hens (Table 1). Cellulose was used (World Minerals, Lompoc, CA) to obtain all diets with 15% crude protein, 2.8 M cal/kg, and 3.5% of total Ca. The phytase was from *Escherichia coli* expressed in *Trichoderma reesei* (Quantum Blue, EC 3.1.3.26, AB Vista, Marlborough, UK) and one FTU was defined as the amount of enzyme required to release 1 mol of inorganic P/min from 0.15 M/dL of sodium phytate at a temperature of 37°C and 5.5 pH.

For each treatment, 12 Bovans White laying hens were placed individually in 40 × 47 cm cages. Hens had ad libitum access to water, with a maximum feed intake of 105 g/hen/day. Facilities were at environmental temperature and hens had a lighting program of 16L: 8D h. All hen handling procedures were approved by the Institutional Subcommittee for the Care and Use of Experimental Animals (SICUAE) of the Veterinary Medicine Faculty of the National Autonomous University of Mexico.

Digestibility test and sampling

Layers were fed for 25 weeks with experimental diets and for additional two weeks with the same diets containing 5 g/kg titanium dioxide (T_iO_2) as an inert marker. For sampling, at age of 65 weeks, all birds were euthanized through intravenous injection of EUTAFIN® (390 mg sodium pentobarbital, 50 mg phenytoin sodium and 1 ml excipients) into the radial vein at a dose of 1 mL/5 kg of BW. The distal ileum sample was collected dissecting the terminal half from the Meckel´s diverticulum portion and remaining up to 2 cm from before the ileo-cecal valve. Digesta content was flushed out with 5 ml of de-ionized water, avoiding pressure to the intestinal mucosa. The digesta from three hens were pooled to one sample, for a total of 52 samples (4 per diet). Samples were frozen at -18°C, and lyophilized for 72 hours.

The gizzard contents were also collected and pooled following the ileal sample order. Contents were homogenized, frozen at -4°C, and lyophilized for 72 hours. Finally, the liver was dissected from hens fed with 0, 300 and 1200 FTU/kg phytase in the diets with 0.67 and 0.87% dLys and from the control diet; they were frozen at -40°C.

Laboratory analysis

Feed: Protein, total lysine, P and Ca contents were analyzed in the basal diet following AOAC (2006) techniques. Exogenous phytase activity in feed samples was analyzed by Enzyme Services and Consultancy (ESC, YstradMynach, UK). Essential and non-essential amino acids, except for tryptophan, were quantified in feed and ileal digesta. Briefly, samples were oxidized using a solution of hydrogen peroxide, formic acid and phenol, and were hydrolyzed with 6M hydrochloric acid for 24 hours at 110°C. Ion exchange chromatography technique was used to separate amino acids, with post-column quantification at 570 nm of ninhydrin (440 nm for proline).

For mineral analysis, samples were incinerated at 500°C for 8 h, and P, Ca, Zn, Cu, Co, Mn, Mg, Fe, K, and Na, and quantified by atomic absorption spectrophotometry technique, following the AOAC (2006b) procedures. The apparent ileal digestibility coefficient (AIDC) of each amino acids and mineral was estimated using the next formulation:

$$AIDC=[(nutrient/TiO_2) \; diet-(nutrient/TiO_2)ileum]/(nutrient/TiO_2) \; diet$$

Low inositol esters: Inositol phosphate esters and inositol were determined by high-performance ion chromatography [12].

Coenzyme Q10: The coenzyme content was determined in pools of three livers, gathered for ileal digesta, giving a total of 28 samples. The extraction of CoQ10 was done following the methodology described for Karadas F, et al. [7], and CoQ10 was quantified as Mattila P and Kumpulainen J [13]. Briefly, the pooled livers were liquefied and mixed with 5% (w/v in H_2O) solution of NaCl (0.7 ml) and ethanol (1 ml) to separate coenzyme in the hexane phase. The extraction was done twice and the hexane phase was combined, evaporated and red is solved in a mixture of methanol/dichloromethane (1:1, v/v). Coenzyme Q10 was analyzed by HPLC using a Hewlett-Packard 1100 equipped with a Vydac 201TP54 column (5 μm, 250 × 4.6 mm, the Separations Goup, U.S.A). The mobile phase was methanol/2-propanol/ethanol (70:15:15, v/v/v), with a flow rate of 0.8 ml/min and the injection volume was 50 μL. Instrumentation was controlled by the HP 3D Chem Station computer program Rev. A. 06.01. The standard employed was Coenzyme Q10 #C9538-1G from Sigma Chemicals Co., U.S.A.

Statistical analysis: Treatments with 0.12% avP were analyzed as factorial arrangement of treatments nested in dietary avP content through Fit Model option of JMP program, categorizing the factors as nominal variables. Further more, avP level was added as source of variation in the ANOVA. Means were separated using the methodology of Tukey or MDS adjusted taking p value as 0.07. The statistical model was:

$$Y=μ+γi+[αm+βn+αβmn]j(i)+ε(ij)k$$

Where μ is the overall mean, γ is i^{th} level of avP level, α is the effect of mth level of dLys, β is the effect of nth dose of phytase and ε is the

experimental error. Regression analysis was applied to continuous values of the factors phytase dose and dLys content, using Fit Model at JMP®program.

Results

There was an interaction between dLys level and phytase dose in digestibility of essential and non-essential amino acids (p<0.05; Table 2 and 3). In the 0.67% dLys level, increasing levels of phytase resulted in a linear positive response in most amino acids, except for Thr. In the 0.77% dLys level, linear positive response was observed for Arg, His, Ile, Lys, and Val, and a quadratic response for Leu and Met, and

Table 1: Composition of basal experimental diet and analyzed nutrient content.

Ingredient	Kilograms
Yellow corn (8%)	690.01
Soybean meal (48%)	199.51
Calcium carbonate	94.31
Salt (NaCl)	4.01
Cellulose	3
Vitamins and minerals[1]	2.41
DL- methionine 84%	1.61
Ortophosphate 18:21	1.31
Yellow pigment 15 g/Kg (*Tagetes erecta*)	1.01
Red pigment 5 g/Kg (*Capsicum annuum*)	0.81
Choline chloride 60%	0.81
BMD-100[2]	0.51
Cyromazine 1%	0.51
Antioxidant[2]	0.21
L- Lysine 76.4%	0.01
Escherichia coli phytase	0
Analyzed nutrient content	
Crude protein (%)[†]	15.1
Metabolizable energy (Mcal/kg)[†]	2.8
Digestible methionine and cysteine (%)[††]	0.65
Digestible methionine (%)[††]	0.38
Digestible lysine (%)[†]	0.67
Digestible threonine (%)[††]	0.61
Total calcium (%)[†]	3.5
Total phosphorus (%)[†]	0.34
Available phosphorus (%)[†]	0.12

[†]Value analyzed; [††]Value calculated

[1]Vitamin premix provided Vitamin A 10,000,000 IU; Vitamin D3 2,500,000 IU; Vitamin E 6,000 IU; Vitamin K 2.5 g; Thymine 1.6 g; Riboflavin 5 g; Cyanocobalamin 0.10 g, Folic acid 0.50 g; Pyridoxine 1.5 g; Calcium pantothenate 10 g; Niacina 30 g; Choline chloride 60% 200 g, Iron 80 g; Manganese 60g; Copper 10 g; Iodine 0.3 g; Zinc 50 g; Selenium 0.30 g; Antioxidant 125 g; Vehicle c.b.p 1,000,000 g per kg diet.

[2]BHA (Butyl hydroxy anisole) 1.2%; BHT (Butyl hidroxy toluene) 9.0%; Ethoxiquine 4.8%; Chelating agents 10

no trend was tracked for Cys and Phe. There was less significance with 0.87% dLys, where Arg, Cys, Ile, and Val exhibited a quadratic tendency, while the digestibility of other essential amino acid was not influenced. In the case of Thr, digestibility was best described by mainly effect of both factors, a linear positive response to dLys level and by quadratic response to phytase dose (Table 2).

About non-essential amino acid digestibility, the response to phytase dose was linear at 0.67% dLys level for Ala, Asp, Glu, Gly, Ser, and Pro. At 0.77% dLys, Pro was the only non-essential amino acid that did not show a linear response. Furthermore, there was a quadratic response to phytase dose in 0.87% dLys, except for Ala and Pro (p<0.05; Table 3). IN ANOVA analyses, level of 0.25% of avP in the diet decreased Asp and His digestibility, while Tyr digestibility was increased (p<0.05; Table 2 and 3).

Digestibility of P was improved with incremental doses of phytase in 0.67 and 0.77% dLys diets (p=0.06). In 0.87% dLys diets, 4800 FTU/kg increased P digestibility regards to no phytase inclusion at both avP content and regards to 300 FTU/kg phytase dose. Furthermore, 1200 FTU/kg showed higher P digestibility than phytase-free 0.12% avP diet.

Mg digestibility increased with phytase addition, particularly in the 0.67% dLys diet, but positive significance was less evident in 0.77% dLys level. Otherwise, it decreased in the 0.87% dLys diet (p<0.05). Increasing dLys content to 0.87% reduced the positive effect of phytase inclusion on K and Na digestibility (p<0.05). Digestibility of Na fit to linear and quadratic responses in diets with 0.67% and 0.77% dLys, respectively. Inclusion of inorganic P of 0.25% avP gave further improvements in digestibility of Ca, Zn, Cu, Fe, Mg, K and Na relative to the group of 0.12% avP diets (p<0.05; Table 4).

Phytase inclusion significantly reduced Ins-6P gizzard concentrations, with statistical differences between the doses 4800 FTU/kg and 300 FTU/kg (p<0.001). 0.25% avP diet had higher Ins-6P amount relative to 0.67% and 0.77% dLys in 0.12% avP diets (p<0.05). Gizzard concentration of Ins-5P in the 0.25% avP diet was higher than the obtained in 0.12% avP phytate-free diet with 0.87% dLys. Doses of 1200 and 4800 FTU/kg phytase gave lower gizzard concentrations of Ins-5P than phytase-free diets with 0.12% avP, while 300 FTU/kg reduced the concentration only to 0.25% avP diet without phytase (p<0.001). Furthermore, gizzard Ins-5P levels in diets with 0.12% avP had less content with 0.67% and 0.77% dLys diets than 0.87% dLys; in addition, this last diet had lower concentrations of Ins-5P than the 0.25% avP diet (p<0.05; table 5).

A significant interaction between phytase doses and dLys levels was observed for gizzard Ins-4P content; the phytase-free diet in 0.87% dLys level gave a higher gizzard content of Ins-4P (p<0.001). Increasing levels of phytase resulted in reductions of the gizzard Ins-4P concentration with decreasing dietary dLys levels (p<0.001; Table 5).

Content of Ins-3P was higher in the phytate-free 0.25% avP diet than in 0.12% avP diets with 1200 and 4800 FTU phytase added. Phytase inclusion at 1200 and 4800 FTU/kg resulted in lower gizzard Ins-3P concentrations compared to 300 FTU/kg in 0.12% avP diets (p<0.001). Diets with 0.77% dLys had lower gizzard contents of Ins-3P than 0.67% and 0.87% dLys diets with 0.12% avP, but not for 0.87% dLys diets with 0.25% avP (p=0.001). For gizzard contents of Ins-2P there was a phytase: dLys interaction, where in 0.67% dLys diets increases of phytase decreased the content of Ins-2P, while in 0.77% and 0.87% diets Ins-2P content increased. Diet with 0.25% avP level gave higher gizzard content of Ins-2P than the average of the diets with

Table 2: Effect of addition of four doses of phytase in three graded levels of digestible lysine on apparent ileal digestibility coefficients of essential amino acids in 67 wk-old White Bovans laying hens.

AvP (%)	Phytase (FTU/kg)	Dig. Lysine (%)	Arg	Cys	His	Ile	Leu	Lys	Met	Phe	Thr	Val
0.12	0	0.67	0.88d	0.69ab	0.86ab	0.83abc	0.86ab	0.84b	0.90ab	0.86abc	0.75ab	0.81ab
	300		0.84e	0.64b	0.80c	0.78c	0.83b	0.77c	0.88b	0.82c	0.67b	0.76b
	1200		0.90abcd	0.73ab	0.86ab	0.86ab	0.88a	0.86ab	0.91ab	0.87abc	0.77a	0.83ab
	4800		0.92a	0.79a	0.90a	0.89a	0.91a	0.90a	0.94a	0.91a	0.82a	0.87a
	0	0.77	0.90abcd	0.73ab	0.87ab	0.85ab	0.88ab	0.89ab	0.91ab	0.88abc	0.77a	0.83ab
	300		0.91abc	0.77a	0.87ab	0.86ab	0.88ab	0.87ab	0.92ab	0.88abc	0.79a	0.84a
	1200		0.92abc	0.78a	0.88ab	0.88ab	0.90a	0.89ab	0.93a	0.90ab	0.80a	0.86a
	4800		0.92a	0.76a	0.89ab	0.88ab	0.90a	0.91a	0.93a	0.88abc	0.80a	0.88ab
	0	0.87	0.89cd	0.71ab	0.87ab	0.83abc	0.86ab	0.88ab	0.91ab	0.83bc	0.74ab	0.81ab
	300		0.92ab	0.76a	0.89ab	0.87ab	0.89a	0.90ab	0.92ab	0.87abc	0.79a	0.85a
	1200		0.91abc	0.77a	0.87ab	0.86ab	0.89a	0.90ab	0.92ab	0.87abc	0.78a	0.85a
	4800		0.90bcd	0.72ab	0.84bc	0.83bc	0.86ab	0.87ab	0.90ab	0.86abc	0.75a	0.81ab
0.25	0		0.89cd	0.75a	0.85bc	0.83ab	0.87ab	0.87ab	0.91ab	0.86abc	0.75ab	0.82ab
		SEM	0.005	0.012	0.006	0.007	0.006	0.007	0.006	0.008	0.010	0.009
Main Effect Means												
0.12	0		0.89bc	0.71b	0.87ab	0.84ab	0.87b	0.87ab	0.91	0.86b	0.76ab	0.82b
	300		0.89c	0.72ab	0.85b	0.84b	0.86b	0.86b	0.91	0.86b	0.75b	0.82b
	1200		0.91ab	0.76a	0.87a	0.86a	0.89a	0.88a	0.92	0.88a	0.78ab	0.85ab
	4800		0.91a	0.76ab	0.87a	0.87a	0.89a	0.89a	0.92	0.88a	0.79a	0.85a
0.25	0		0.90abc	0.75ab	0.85b	0.83ab	0.87ab	0.87a	0.91	0.86ab	0.75ab	0.82ab
0.12		0.67	0.89bL	0.71bL	0.85bL	0.84bL	0.87bL	0.84bL	0.90bL	0.87abL	0.75b	0.82bL
		0.77	0.91aL	0.76a	0.88aL	0.87aL	0.89aQ	0.89aL	0.92aQ	0.88a	0.79a	0.85aL
		0.87	0.92aQ	0.74abQ	0.87ab	0.85abQ	0.87ab	0.89a	0.91ab	0.86b	0.77ab	0.83bQ
0.25			0.89ab	0.75ab	0.85b	0.83ab	0.87ab	0.87ab	0.91ab	0.86ab	0.75ab	0.82ab
0.12			0.90	0.74	0.87b	0.85	0.88	0.87	0.91	0.87	0.77	0.83b
0.25			0.89	0.75	0.85a	0.83	0.87	0.87	0.91	0.86	0.75	0.82a
p-values												
	Phytase		0.001	0.011*	0.048*	0.004**	0.002**	<.001	0.088	0.019*	0.009**Q	0.004**
	Lysine		0.001	0.008**	0.008**	0.010*	0.017*	<.001	0.037*	0.026*	0.007**L	0.009**
	Lysine*Phytase		<0.001	0.006**	<0.001	<0.001	0.001	<.001	0.008*	0.018*	<0.001	0.001
	AvP		0.353	0.620	0.063*	0.112	0.513	0.974	0.936	0.666	0.323	0.582

0.12% avP (p<0.001; Table 5). Layers fed with 1200 FTU/kg had higher gizzard digest a levels of inositol than those fed with phytase-free diets with 0.12% and 0.25% avP levels. Additionally, 0.25% avP diet had lower gizzard inositol content than 0.12% avP diets (p<0.05; Table 5).

In 0.67% dLys, there were no differences between phytase doses in CoQ10 content, but in 0.87% phytase-free diet resulted in higher CoQ10 liver content than 300 and 1200 FTU/kg doses. Including 1200 FTU/kg in 0.87% dLys level reduced the CoQ10 concentration compare to 0, 300 and 1200 FTU/kg phytase addition (p<0.001). Furthermore, 0.25% avP diet gave lower liver content than the average of the 0.12% avP diets (p=0.001; Table 5).

Discussion and Conclusion

In layers previous studies regard to amino acid digestibility effect of phytase are few and results are generally inconclusive. While Jalal MA, et al. [14] observed effect on the digestibility of Met, Cys, Ala,

Table 3: Effect of addition of four doses of phytase in three graded levels of digestible lysine on apparent ileal digestibility of non-essential amino acids in 67 wk-old White Bovans laying hens.

AvP (%)	Phytase (FTU/kg)	Diges. Lysine (%)	Ala	Asp	Glu	Gly	Pro	Ser	Tyr
0.12	0	0.67	C 0.83[abcd]	0.82[abc]	0.87[ab]	0.78[ab]	0.83[ab]	0.81[ab]	0.85[ab]
	300		0.79[d]	0.77[c]	0.85[b]	0.71[b]	0.80[b]	0.75[b]	0.82[ab]
	1200		C 0.85[abcd]	0.84[ab]	0.89[ab]	0.79[ab]	0.85[ab]	0.83[a]	0.84[ab]
	4800		0.90[a]	0.88[a]	0.92[a]	0.84[ab]	0.89[a]	0.86[a]	0.89[a]
	0	0.77	0.84[abcd]	0.84[ab]	0.88[ab]	0.80[a]	0.85[ab]	0.83[a]	0.86[ab]
	300		0.84[abcd]	0.85[ab]	0.90[ab]	0.80[a]	0.87[a]	0.84[a]	0.87[a]
	1200		0.87[abc]	0.86[ab]	0.91[a]	0.83[a]	0.88[a]	0.86[a]	0.87[a]
	4800		0.88[ab]	0.87[ab]	0.91[a]	0.84[a]	0.88[a]	0.86[a]	0.84[ab]
	0	0.87	0.81[cd]	0.82[abc]	0.87[ab]	0.77[ab]	0.83[ab]	0.81[ab]	0.80[b]
	300		0.86[abcd]	0.86[ab]	0.90[a]	0.82[a]	0.86[ab]	0.84[a]	0.83[ab]
	1200		0.86[abcd]	0.85[ab]	0.90[a]	0.81[a]	0.87[a]	0.85[a]	0.87[ab]
	4800		0.82[bcd]	0.82[abc]	0.88[ab]	0.77[a]	0.85[ab]	0.82[a]	0.86[ab]
0.25	0		0.85[abcd]	0.81[bc]	0.87[ab]	0.78[ab]	0.85[ab]	0.82[a]	0.89[a]
	Sem		0.008	0.007	0.006	0.010	0.008	0.008	0.008
Main Effect Means									
0.12	0		0.83[c]	0.83[ab]	0.88[b]	0.78[ab]	0.84[b]	0.81[bc]	0.84[b]
	300		0.83[bc]	0.83[ab]	0.90[ab]	0.78[b]	0.84[ab]	0.81[c]	0.84[b]
	1200		0.86[ab]	0.85[a]	0.89[ab]	0.81[ab]	0.87[ab]	0.84[ab]	0.86[ab]
	4800		0.87[a]	0.86[a]	0.87[a]	0.82[a]	0.87[a]	0.85[a]	0.86[ab]
0.25	0		0.85[abc]	0.81[b]	0.87[a]	0.78[ab]	0.85[ab]	0.82[abc]	0.89[a]
0.12		0.67	0.84[ab]**L**	0.83[b]**L**	0.88[b]**L**	0.78[b]**L**	0.84[ab]**L**	0.81[b] **L**	0.85[b]
		0.77	0.86[a]**L**	0.86[a]**L**	0.90[a]**L**	0.82[a]**L**	0.87[ab]**Q**	0.84[a] **L**	0.86[ab]
		0.87	0.84[b]	0.84[ab]**Q**	0.90[ab]**Q**	0.79[ab]**Q**	0.85[a]	0.83[ab]**Q**	0.84[ab]
0.25			0.85[ab]	0.81[b]	0.87[ab]	0.78[ab]	0.85[b]	0.82[ab]	0.89[a]
0.12			0.85	0.84[a]	0.89	0.80	0.85	0.83	0.85[b]
0.25			0.85	0.81[b]	0.87	0.78	0.85	0.82	0.89[a]
P-values									
Phytase			0.001***	0.007**	0.003***	0.012*	0.004***	0.001***	0.059*
Lysine			0.069*	0.011*	0.025*	0.019*	0.020*	0.003***	0.063*
Lysine*Phytase			0.003**	0.0003***	0.005***	0.001***	0.049*	0.001***	0.005**
AvP			0.961	0.028*	0.128	0.406	0.633	0.332	0.008**

and Glu with 250 and 300 FTU/kg phytase in a corn and soybean meal based diet, Agbede JH, et al. [10] did not find any positive effect adding 1500 FTU/kg phytase to sunflower and rapeseed meal testing diets with two levels of non-phytic P (8.0 and 4.0 g/kg). In the present study, the sampling region, the sampling method, and the marker used can partly explain the significance on amino acid digestibility. Thus, Rezvani M, et al. [15] recommended taking samples from middle and terminal sections of the ileum (between Meckel's diverticulum and ileo-caecal-colonic junction), due to almost all amino acids have been absorbed by this stage and microbial activity is low. Adedokun SA, et al. [16] advised flushing out the digesta with deionized water rather than squeezing, to reduce contamination with endogenous amino acids, meanwhile the use of titanium dioxide lead to less variability.

Ravindran V, et al. [17] reported linear increases in amino acid digestibility with increasing phytase dose when a dLys deficient diet (1.00%) was tested in chicks. The current experiment intended to demonstrate an increase in Lys supply by phytase, however Lys release from enzyme was not enough to compensate the reduction in dietary dLys from 0.67% to 0.77%, or from 0.77 to 0.87%.

Selle PH, et al. [18] reported that addition of 500 FTU/kg phytase improved performance in 7-28 day-old broilers fed a dLys deficient (10.0 g/kg) diet, but not in a dLys adequate (11.8 g/kg) diet. Consequently, an interaction between phytase and Lys contents was observed on the digestibility of Arg, Lys, Phe, Asp, Glu, Gly and Ser. These authors explained the mainly Lys effect based on the stimulus of epithelia transporters, such as the b^0, + system at intestinal level for amino acids. Nevertheless, since Lys is the reference amino acid for ideal amino acid profile, its availability can strongly influence others essential amino acids for protein, and for that in the present Lys significantly affect amino acids avalilability.

Table 4: Effect of addition of four doses of phytase in three graded levels of digestible lysine on apparent ileal retention of minerals in 67 wk-old White Bovans laying hens.

AvP (%)	Phytase (FTU/kg)	Diges. Lysine (%)	P	Ca	Zn	Fe	Mg	Mn	Cu	K	Na
0.12	0	0.67	0.44[b]	0.37	0.25	0.15	0.43[abc]	0.06	0.49	0.79[ab]	-0.25[cd]
	300		0.44[b]	0.52	0.35	0.05	0.36[bcd]	-0.03	0.29	0.72[ab]	-0.53[d]
	1200		0.57[ab]	0.35	0.09	0.34	0.42[abc]	-0.18	0.09	0.78[ab]	-0.02[abc]
	4800		0.73[a]	0.51	0.22	0.16	0.45[ab]	0.04	0.55	0.86[a]	0.10[abc]
	0	0.77	0.45[b]	0.40	0.24	0.21	0.41[abcd]	0.03	0.50	0.78[ab]	-0.19[cd]
	300		0.60[ab]	0.66	0.36	0.37	0.48[a]	0.20	0.40	0.82[ab]	-0.23[abc]
	1200		0.71[a]	0.65	0.39	0.25	0.45[ab]	0.10	0.32	0.84[ab]	0.27[ab]
	4800		0.75[a]	0.68	0.41	0.21	0.46[ab]	0.05	0.44	0.84[ab]	0.05[abc]
											5
	0	0.87	0.59[ab]	0.47	0.41	0.26	0.51[a]	0.16	0.42	0.81[ab]	0.15[abc]
	300		0.72[a]	0.31	0.20	0.05	0.47[ab]	-0.03	0.19	0.82[ab]	0.05[cd]
	1200		0.71[a]	0.62	0.44	0.31	0.41[abcd]	0.16	0.38	0.82[ab]	0.11[abc]
	4800		0.73[a]	0.61	0.39	0.27	0.32[cd]	0.23	0.29	0.75[ab]	-0.09[bc]
0.25	0		0.56[ab]	0.73	0.50	0.48	0.31[d]	0.19	0.53	0.72[b]	0.39[a]
	SEM		0.027	0.063	0.052	0.051	0.024	0.066	0.036	0.015	0.049
Main Effect Means											
0.12	0		0.50[c]	0.42	0.30	0.21	0.45	0.09	0.47[a]	0.79	-0.02[cd]
	300		0.59[bc]	0.50	0.30	0.16	0.44	0.05	0.19[b]	0.79	-0.24[d]
	1200		0.66[ab]	0.54	0.30	0.30	0.43	0.03	0.37[a]	0.81	0.12[b]
	4800		0.74[a]	0.60	0.34	0.22	0.41	0.11	0.43[a]	0.81	0.02[bc]
0.25	0		0.56[bc]	0.73	0.50	0.48	0.31	0.19	0.53[a]	0.72	0.39[a]
0.12		0.67	0.55[b] **L**	0.44	0.23[b]	0.18	0.42	-0.03	0.36	0.78	-0.17[c]**L**
		0.77	0.63[ab]**Q**	0.60	0.35[ab]	0.26	0.45	0.10	0.41	0.82	0.040[b]**Q**
		0.87	0.69[a] **Q**	0.50	0.36[a]	0.22	0.43	0.13	0.32	0.80	-0.02[b]
0.25			0.56[ab]	0.73	0.50[a]	0.48	0.31	0.19	0.53	0.72	0.39[a]
0.12			0.62	0.51[b]	0.31[b]	0.22[b]	0.43[a]	0.07	0.36[b]	0.80[a]	-0.05[b]
0.25			0.56	0.73[a]	0.50[a]	0.48[a]	0.31[b]	0.19	0.53[a]	0.72[b]	0.39[a]
p-values											
Phytase			<.0001	0.204	0.951	0.277	0.579 **Q**	0.844	<.0001	0.483 **L**	<.0001
Lysine			0.001***	0.131	0.079	0.401	0.503	0.129	0.145	0.155	0.002**
Lysine*Phytase			0.062*	0.393	0.131	0.201	0.035*	0.452	0.093	0.008**	0.001***
AvP			0.226	0.059*	0.048*	0.006**	0.006**	0.298	0.011*	0.003**	<.0001

Hydrolyses of Phyt-ac was reported to increase by 50% and 66%, respectively, adding 250 and 500 FTU/kg of a 3-phytase to laying hen diets [19], meanwhile in another study, ileal P digestibility showed improvements of 39.0%, 44.6%, and 51.3% with 150, 300, and 450 FTU/kg 3-phytase, respectively [20]. In our study, increasing levels of phytase improved the release of P from diets, but higher doses were required to release higher levels of phytate-P. On the other hand, Ca digestibility was not influenced by phytase, notwithstanding of modifications in P digestibility influence Ca absorption, for maintaining the optimal Ca:P ratio in biological process such as egg shell formation.

No consistent effects of phytase were observed in this study on Zn, Cu, Co, Mn, Mg, and Fe digestibility, in spite of chelation process of these minerals by Phyt-ac [21], likely because contents in diets was covering the mineral requirements of the hens, disguising phytase improvements on extra availability. In the case of Na, intestinal mucus production increases in the presence of phytate, which can result in higher losses of mucus component, including Na [22]. When an exogenous phytase is added, a direct effect on Na homeostasis is expected, such as was observed in the present study. Cowieson A, et al. [4] reported that including 1000 FTU/kg phytase offset the

Table 5: Effect of addition of four doses of phytase in three graded levels of digestible lysine on content of hydrolyses molecules of phytic acid in gizzard digesta and coenzyme Q10a in liver of 65 week-old White Bovans laying hens.

t	Phytase (FTU/kg)	Diges. Lysine (%)	Ins6P[c]	Ins5P	Ins4P	Ins3P	Ins2P	Inositol	CoQ10[b]
			N mol/g Dry Weight						
0.12	0	0.67	4159	400	3[d]	557	167[d]	656	392.9[b]
	300		2108	259	93[cd]	554	546[b]	871	338.8[bc]
	1200		1859	205	45[cd]	383	394[bcd]	103	380.2[b]
	4800		489	43	15[d]	328	207[cd]	629	---
	0	0.77	2991	362	49[cd]	411	321[bcd]	293	---
	300		1627	192	83[cd]	380	373[bcd]	450	---
	1200		701	78	34[cd]	318	493[bc]	78	---
	4800		326	43	5[d]	303	452[bcd]	841	---
	0	0.87	3260	640	769[a]	648	921[a]	441	607.5[a]
	300		2424	428	334[b]	511	911[a]	572	c300.5[bcd]
	1200		1511	252	260[bc]	376	963[a]	1078	217.5[d]
	4800		1480	283	123[bcd]	350	1034[a]	1145	---
0.25	0		4187	827	192[bcd]	364	882[a]	119	257.0[cd]
		SEM	253.5	44.07	26.80	26.70	37.27	103.5	25.51
			Main Effect Means						
0.12	0		3470[a]	467[b]	273[a]	539[a]	470[b]	463[bc]	500.2[a]
	300		2053[b]	293[bc]	170[ab]	482[ab]	610[b]	631[abc]	319.7[b]
	1200		1357[bc]	178[c]	113[bc]	359[c]	617[b]	963[a]	298.8[b]
	4800		765[c]	123[c]	48[c]	327[c]	564[b]	871[ab]	---
0.25	0		4187[a]	827[a]	192[abc]	364[bc]	882[a]	119[c]	257.0[b]
0.12		0.67	2154[b]	227[c]	39[c]	456[a]	329[b]	797	370.6
		0.77	1411[b]	169[c]	43[c]	353[b]	410[b]	591	---
		0.87	2169[b]	401[b]	372[a]	471[a]	957[a]	809	375.1
0.25			4187[a]	827[a]	192[b]	364[ab]	882[a]	119	257.0
0.12			1911[b]	265[b]	151	427	565[b]	732[a]	372.9[a]
0.25			4187[a]	827[a]	192	364	882[a]	119[b]	257.0[b]
			p-values						
Phytase			<.0001 **Q**	<.0001 **Q**	<.0001 **L**	<.0001 **Q**	0.025*	0.005** **Q**	<.0001 **L**
Lysine			0.025*	0.001	<.0001 **L**	0.001	<.0001 **L**	0.156	0.831
Lysine*Phytase			0.469	0.879	<.0001	0.289	0.011*	0.314	<.0001
AvP			<.0001	<.0001	0.392	0.191	<.0001	0.002**	0.001

[a]Content of coenzyme Q10 is reported in three phytase doses and two levels of digestible lysine
[b]Coenzyme Q10
[c]Low inositol ester
Statistically significant *p ≤ 0.05, **p ≤ 0.01, ***p ≤ 0.001. a-b Show significant differences among treatment means.

negative impact of Phyt-ac on Na excretion in broilers; whereas, they did not observe any changes in the excretion of Zn, Fe, Mg, Mn, and Cu. Moreover, the likely influence on Na/K ATPase intestinal pump by changes in Na and K digestibility, could affect general nutrient absorption, other than P, as was obtained.

In poultry, Phyt-ac degradation increases with increasing phytase dose, with a linear positive relation observed in broilers and also in laying hens [9]. Persson H, et al. [21] reported a lower binding strength for Cu, Zn and Cd when Ins-4 and 3P were tested compare to Ins-5 and 6P. Additionally, Xu P, et al. [23] observed *in vitro* that Ca and Zn solubilities were increased when Phyt-ac was degraded to Ins-3P. In the current study, 300 FTU/kg, which is generally the standard dose of phytase, was efficient in the initial breakdown of Phyt-ac, meanwhile 1200 FTU/kg increased free inositol release. Thus, high phytase doses

increased the presence of Ins-P breakdown products with weaker covalent binding strength, and then the availability of nutrients that are susceptible to chelation by Phyt-ac.

One important result to highlight in this study was the effect of dLys on Phyt-ac degradation by phytase; this can be explained following the electrostatic interactions between molecules. Kaup SM and Greger J [24] investigated the impact chloride salt type on intestine absorption, bone deposition, and excretion of P, Ca and Mg in rats, concluding that Lys chloride was able to increase Ca excretion. Also, when Lys chloride and Ca were supplied in the diet, P and Mg absorption in the intestine and Mg deposition in bone were affected negatively. Thus, anions such as chloride can influence the electrostatic interactions among dietary molecules, depending of type of Ins-P present, and can result in deeply changes in digestive process.

Therefore, in the present study Lys chloride addition into the diet could have altered the solubility and hydrolysis of Phyt-ac, resulting in mainly effects of dLys level into the diets. Similarly, Banks K, et al. reported linear reductions in P retention with increasing dietary Cu concentration from 125 to 375 ppm, but not for 10 and 62.5 ppm [25]. Furthermore, Wilcock P and Walk CL [26] reported that Cu form included in diet influences the solubility of Cu-phytate complexes, since sulfate, chloride and tribasic chloride Cu resulted in higher levels of insoluble Cu than citrate and lysine sources at pH 5.5. This reinforces the idea that molecular composition highly influences Phyt-ac stability and susceptibility to phytase action.

Laying hens produce phosphatases able to hydrolyze Phyt-ac up to 35% to releasing inositol and free phosphate groups [27]. Truong HH, et al. [28] reported 63% overall hydrolysis of phytate in broilers, with 24% corresponded to endogenous enzymes action and 39% to 1000 FTU/kg for an exogenous phytase. Recently, Wilcock P and Walk CL [26] established that using 1500 FTU/kg phytase, a high dose in poultry meat production, more than 85% of Phyt-ac is hydrolyzed, meanwhile the standard dose of 500 FTU/kg degrades 29% or 34%. In our study, inclusion of 4800, 1200 and 300 FTU/kg phytase reduced Ins-P content by 78%, 61%, and 41%, respectively, in gizzard digesta.

Supplementation with 0.13% inorganic P was not equivalent in P release by the phytase doses evaluated in the current trial. Liu N, et al. [29] found higher P digestibility using 0.28% avP rather than including 300 FTU/kg from three different commercial phytase in Hy-line Brown layers fed 0.58% dLys. By contrast, in the present study 300 FTU/kg phytase gave a higher P digestibility (59%) than inorganic-P inclusion (56%), and while P was more available with increasing phytase dose, the effect was lower in diets with 0.87% dLys. Furthermore, the addition of 0.13% P as monocalcium phosphate increased the presence of Ins-6 and 5P in gizzard digesta and reduced Ins-2P and inositol concentrations. Shastak Y, et al. [30] found less Ins-6P hydrolysis in ileum digesta and excreta of 20-24 day-old broilers with increasing inorganic-P from 0.075% to 0.15% using monocalcium phosphate monohydrate. Similarly, Li W, et al. [31] observed an increase of Ins-6P presence in proventriculus and gizzard digesta in 13 day old chickens with 0.45% non-phytate P relative to 0.28% non-phytate P when fed a mixed basal diet with monocalcium phosphate as inorganic P source. Inorganic P content is one of the most important negative factors in phytase hydrolysis; hence, the low inorganic-P level in the current trial (0.12%) allowed phytase to have a significant effect on amino acid digestibility, as previously reported with 0, 250 and 500 FTU/kg for amino acid and P digestibility [19].

High concentrations of cell CoQ10 content is indicative of oxidative status, mainly due to catabolism of lipids, carbohydrates and proteins [8]. In the diet with 0.87% dLys without phytase, more catabolism of extra amino acid was presented altering the citric acid cycle where CoQ10 is a key molecule. Conversely, CoQ10 content showed less variation to phytase in 0.67% dLys because the enzyme can regulate the requirements of P but not the lysine. In the 0.87% dLys diet, inorganic P supplementation and phytase addition decreased liver CoQ10 concentration by 58%, although the greatest effect was observed with 1200 FTU/kg addition.

Karadas F, et al. (2010) observed a higher liver content of CoQ10 and better performance in broilers fed diets with 500 FTU/kg (99.95 μg/g) and 12500 FTU/kg (119.7 ug/g) of phytase compared to 250 FTU/kg (63.86 ug/g) [7]. Additionally, Karadas F, et al. [32] reported no change in the coenzyme concentration using 12500 FTU/kg or P inorganic supplementation (129.9 and 119.7 ug/g, respectively). The above-mentioned author assumed that high CoQ10 enhances oxidative status of cellular membranes in the liver, and it allows better productive performance. In our case, opposite result was obtained, and it is likely due to differences in intrinsic physiologically characteristics of layer as age, but no previous report are available to confirm the current results. The only available research quantifying CoQ10 content reported improvement of 8% when a source of CoQ10 was supplemented at diet [33].

In conclusion, the increase of phytase dose has improved the availability of essential, non-essential amino acids, P and Na, with less effect on high dLys diet. Moreover, higher inclusion promotes the presence of lower inositol esters (Ins5-2P) and inositol from Phyt-ac, and affect CoQ10 content in liver. The response to inorganic P addition contrasted to that of standard dose of phytase (300 FTU/kg), even though for P digestibility.

Acknowledgments

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