

## Effects of exogenous tannase enzyme on growth performance, antioxidant status, immune response, gut morphology and intestinal microflora of chicks fed grape pomace

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### Abstract

An experiment was conducted to study the effects of dietary addition of tannase to feed of chicks including grape pomace (GP) on growth performance, antioxidant status, immune response, blood parameters, gut morphology, intestinal microflora, liver function, and histopathological responses. The experimental diets were i) control (corn and soybean diet) (C); ii) C+10%GP; iii) C+10%GP+T1 (500 mg/kg tannase enzyme); and iv) C+10% GP+T2 (1000 mg/kg tannase enzyme). At 10 days old, the bodyweight (BW) and average daily gain (ADG) of the birds fed the diet supplemented with GP was lower compared with the control group. In contrast with the control, supplementation of diets with 10%GP+1000 mg/kg tannase elevated superoxide dismutase (SOD) and glutathione peroxidase (GSH-Px) activities and depressed the malondialdehyde (MDA) level in serum. The addition of GP to the chicken diets had a significant impact on the total anti- sheep red blood cells (SRBC) titers and IgG, and IgM antibodies at 21 and 42 days old. The muscularis thickness of the chicken duodenum decreased in the Trial 3 compared with control groups. The inclusion of GP in the chicken diets reduced the concentration of *Escherichia coli* and increased that of *Lactobacillus* compared with the control. The results of the present study suggest that the inclusion of up to 10 percent GP in diets did not adversely affect broiler growth performance, and supplementation of tannase improved the antioxidant status and immune responses, and increased the caecal populations of beneficial bacteria in the cecum of broiler chickens.

**Keywords:** Broiler chickens, histopathological responses, glutathione peroxidase, polyphenol, sheep red blood cells

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### Introduction

Grapes (*Vitis vinifera*) are one of the world's largest fruit crops, with an annual production of 77 million metric tons (FAO, 2013). Large quantities of residues are generated in processing grapes. These residues cause problems in economic and ecological terms. Thus, any useful application for these by-products could represent an interesting advance in the maintenance of environmental equilibrium and also an economic evaluation of the raw material (Abarghuei *et al.*, 2010). Grape pomace (GP) is a grape by-product consisting of pressed seeds, skins and stems, and is a rich source of flavonoids, including monomeric phenolic compounds, such as (+)-catechins, (-)-epicatechin, and (-)-epicatechin-3-gallate and dimeric and oligomeric proanthocyanidins (Goni *et al.*, 2007). These polyphenols were considered anti-nutritional factors because their presence in certain ingredients, such as legumes, sunflowers and sorghum, had negative effects on animal nutrition. Probably one of the best-known properties of polyphenolic compounds is their capacity to bind and precipitate proteins. This protein binding capacity is common to most polyphenols due to their high degree of hydroxylation. Low molecular weight phenols are, however, not able to precipitate proteins. Oligomers must contain at least three flavonol subunits to precipitate protein effectively. Additionally, these compounds have the capacity to act as powerful antioxidants by scavenging free radicals and terminating oxidative reactions (Yilmaz & Toledo, 2004). Various plant polyphenolic compounds have been studied widely because of their antioxidant and antimicrobial activities. Plant extracts that are rich in these polyphenols have been used successfully to protect against diseases caused by organisms such as

*Staphylococcus aureus*, *Escherichia coli*, *Candida albicans*, and *Campylobacter* (Tepe *et al.*, 2004). Therefore, GP polyphenols in grape by-products could be used effectively as feed supplements to improve the antioxidant status and immunity of birds. However, the use of such natural antioxidants in animal nutrition could be limited owing to the low bioavailability of grape polyphenols, and might be improved with exogenous enzymes.

Enzyme supplementation is a technique that has had increasing applicability for improving the nutritional characteristics of by-products and is widely used in animal nutrition. A recent *in vitro* study (Chamorro *et al.*, 2012) reported that the addition of tannase released polyphenols entrapped in GP, increasing its antioxidant activity. Tannase catalyses the hydrolysis of carboxylic ester bonds in the molecules of hydrolysable tannins and gallic acid esters (Lekha & Lonsane, 1997). Tannase is produced by certain filamentous fungi, mainly *Aspergillus*, *Penicillium*, *Fusarium* and *Trichoderma* (Libuchi & Monida Yand Yamada, 1967; Rajakumar & Nandy, 1983; Kawakubo *et al.*, 1991; Lekha & Lonsane, 1994; Bajpai & Patil, 1996; Garcia-Pena *et al.*, 1999) and bacteria (Deschamps *et al.*, 1983; Skene & Brooker, 1995). Yeasts (Aoki *et al.*, 1976) also produce this enzyme. Most of the work related to tannase has considered production, extraction, purification, and characterization aspects of the enzyme obtained by microbiological methods, using filamentous fungi, mainly *Aspergillus*, through processes of submerged culture. In general form, fungal tannase is produced by submerged cultures (Lekha & Lonsane, 1994). It is the chemical structure of polyphenols, rather than the concentration, that determines the rate and extent of absorption and the nature of the metabolites circulating in the plasma. The hydrolysis of complex polyphenols into more digestible phenols might increase the amount of active substances that could be metabolized easily, improving its nutritional value and rendering this by-product more suitable for use as an animal ingredient.

The main objective of this experiment was to study the effects of dietary addition of tannase to the feed of chicks including GP on growth performance, antioxidant status, immune response, blood parameters, gut morphology, intestinal microflora, liver function and histopathological responses.

## Materials and Methods

Red grape pomace (*Vitis vinifera*) was obtained from TATAO Corporation (Urmia-West Azerbaijan, Iran). The proximate composition of GP was analysed according to procedures described by the Association of Official Analytical Chemists (AOAC, 2000) (Table 1). Total polyphenol content in GP, diet, ileal digesta and excreta were determined after extraction with methanol/HCl 99/1 (v/v) as an extraction solvent using the Folin-Ciocalteu colorimetric method (Singleton & Rossi, 1965). Gallic acid was employed as a calibration standard and results were expressed as gallic acid equivalents (mg gallic acid/g of dried samples). The GP was used as a source of dietary fibre and polyphenols in the chicken diets.

**Table 1** Proximate composition of grape pomace and total polyphenol content in grape pomace, control and 10 % grape pomace diets<sup>1</sup>

Parameters	Concentration	GP and diets	Total polyphenols (mg gallic acid equivalent /gDM)
Dry matter (g/kg)	914.5 ± 0.37	GP	33.92 ± 0.48
Protein (g/kg)	89.4 ± 0.16	Control diet	3.66 ± 0.35
Fat (g/kg)	70 ± 0.91	10 % GP diet	11.34 ± 0.64
Ash (g/kg)	32.6 ± 0.43		
Fibre (g/kg)	302 ± 1.97		
Ca (g/kg)	5.2 ± 0.09		
P (g/kg)	2.9 ± 0.08		
Gross energy (kcal/kg)	4397.63 ± 164.22		

<sup>1</sup>Data are the mean of three determinations ± SD

GP: grape pomace

A total of 200 day-old male broiler chicks (Ross 308) were purchased from a local hatchery. On arrival, all birds were weighed (average bodyweight  $40 \pm 2$  g) and randomly assigned to one of four treatments, with five replicate pens/treatment and ten chickens/pen. Dietary treatments were formulated to meet or exceed the nutrient requirements of broilers provided by Ross Broiler Manual (Aviagen, 2014) (Table 2). Experimental diets consisted of i) control: corn, soybean diet (C); ii) C+10% of GP; iii) C+10% GP+T1 (500 mg/kg tannase enzyme); and iv) C+10% GP+T2 (1000 mg/kg tannase enzyme). The straw was replaced with GP in the experimental diets. The feeding regimen consisted of starter (0 to 10 days), grower (11 to 24 days), and finisher diets (25 to 42 days). Chickens were raised in floor pens (100 x 120 cm) and had free access to feed and water for the entire experimental period (days 0 – 42). The room temperature gradually decreased from 33 to 22°C on day 28 and then remained constant. The lighting programme consisted of 20 hours of light and 4 hours of darkness.

All experimental procedures were evaluated and approved by the Institutional Animal Care and Ethics Committee of Mohaghegh Ardabili University. An enzyme with tannase activity (T) was used. It was supplied by Kikkoman Food Products Co (Edogawa Plant, Japan) and contained tannin acylhydrolase (500 U/g, EC 3.1.20).

**Table 2** Ingredients and nutrient composition of basal diets

Ingredients	Starter (days 0–10)	Grower (days 11–24)	Finisher (days 25–42)
Corn	42.53	44.52	48.75
Soybean meal (44% CP)	41.18	37.87	32.76
Soybean oil	5.99	7.27	7.77
Straw <sup>4</sup>	5.93	6.40	7.06
Grape pomace <sup>3</sup>	0	0	0
Dicalcium phosphate	1.75	1.57	1.40
Limestone	1.10	0.99	0.94
Sodium chloride	0.35	0.35	0.35
Vitamin premix <sup>1</sup>	0.25	0.25	0.25
Mineral premix <sup>2</sup>	0.25	0.25	0.25
DL-Methionin	0.36	0.32	0.29
L-Lysine	0.22	0.15	0.14
L-Threonine	0.10	0.07	0.05
Calculated analysis			
ME	2900	3000	3100
Crude protein	22.22	20.81	18.89
Met + cystine	1.04	0.96	0.88
Lysine	1.39	1.25	1.12
Ether extract	7.87	9.19	9.80
Crude fibre	6.20	6.20	6.20
Ca	0.93	0.84	0.77
Available P	0.46	0.42	0.38

<sup>1,2</sup>Each kg of vitamin and trace mineral premix provided vitamin A: 900 IU; vitamin D: 2000 IU; vitamin E: 18 IU; vitamin K: 2 mg; vitamin B1: 1.8 mg; vitamin B2: 6.6 mg; vitamin B6: 3 mg; vitamin B12: 15 µg; niacin: 30 mg; pantothenic acid: 10 mg; biotin: 0.1 mg; folic acid, 1.25 mg; choline chloride: 200 mg; Fe: 50 mg; Cu: 10mg; Mn: 100 mg; Zn: 85 mg; I: 0.8 mg; Se: 0.2 mg

<sup>3</sup>Experimental diets consisted of i) control corn, soybean diet (C); ii) C+10% GP; iii) C+10% GP+T1 (500 mg/kg tannase enzyme); and iv) C + 10% GP+T2 (1000 mg/kg tannase enzyme)

<sup>4</sup>The straw was replaced with grape pomace in the experimental diets; CP : crude protein; ME: metabolizable energy.

Birds and feed were weighed at 10, 24, and 42 days old. The values of average daily feed intake (ADFI) and ADG were recorded at various periods, and the feed-to-gain ratio was calculated. Mortality was

recorded as it occurred. However, ADFI and F/G were corrected for the mortality of related groups. The economic evaluation was made in terms of feed cost per kg live weight gain during the differentiation and overall study period. At the end of the experiment (on day 42), after overnight fasting, two birds from each pen with bodyweight close to the pen mean were selected and slaughtered. Viscera were removed manually, and carcass characteristics (carcass yield, breast, thigh and abdominal fat) were determined. Then, internal organ weights were measured. All carcass data are presented based on percentage of live weight of each bird.

At 42 days old, two blood samples were collected from two birds/replicate following approximately eight hours fasting to determine serum biochemical parameters and antioxidative status. The first samples were collected into vials containing EDTA and centrifuged for 10 min at 3000xg, then plasma was collected and stored at -20°C until analysis. The concentrations of total protein, glucose, uric acid, triglycerides (TG), total cholesterol (CHOL), high-density lipoprotein cholesterol (HDL-C), low-density lipoprotein cholesterol (LDL-C), aspartate aminotransferase (AST) and alanine aminotransferase (ALT) in plasma samples were analysed with an automatic biochemical analyser (HITACHI912. Japan), following the instructions of the corresponding reagent kit (Audit Diagnostics Co. Ireland). The second blood sample was collected in a heparinized graduated centrifuge tube to obtain hemolysate. This was used to evaluate antioxidative status by measuring the malondialdehyde (MDA), superoxide dismutase (SOD) and glutathione peroxidase (GSH-Px) activity. GSH-Px activity was determined with a commercial enzyme kit (Ransel, RANDOX/RS-504 supplied by Randox Laboratories, Crumlin, UK). The SOD activity was determined with a commercial enzyme kit (Ransod, RANDOX/SD-125 supplied by Randox Laboratories). The MDA was analysed with a commercial kit (Biocore Diagnostik Co. Germany) and MDA was measured as a decomposed product of lipid peroxidation with 2-TBA using the colorimetric method with a spectrophotometer at a wavelength of 532 nm (Placer *et al.*, 1966). At nine days old, Newcastle and influenza antigens were injected subcutaneously with 0.2 ml per chick with a dual vaccine of Newcastle and influenza. Also, the chicks were orally vaccinated against Newcastle disease at 19 days old. To assay the primary and secondary antibody responses against sheep red blood cell (SRBC), at 14 and 35 days old, two birds/replicate were immunized intramuscularly with 0.25 ml 10% SRBC in phosphate-buffered saline (Leshchinsky & Klasing, 2001). On days 21 and 42, the same birds were bled to determine antibody titers against SRBC and Newcastle disease virus. Subsequently, antibody titers against this virus were measured separately by haemagglutination inhibition (HI). The HI antibodies were then converted to log<sub>2</sub>. The serum from each sample was collected, heat inactivated at 56°C for 30 min, and then analysed for total IgM and IgG (mercaptoethanol-resistant) anti-SRBC antibodies by the microhaemagglutination technique described by Qureshi & Havanstein (1994).

Cellular immunity was assessed by a cutaneous basophil hypersensitivity test *in vivo* with phytohemagglutinin. At day 38, the toe web between the third and fourth digits of the right foot was measured in millimetres with a constant tension micrometer. Immediately after measurement, 100 µg phytohemagglutinin, suspended in 0.10 ml sterile saline, was injected into the toe web. The toe web swelling was measured 24 and 48 hours after injection. The response was determined by subtracting the skin thickness of the first measurement from that of the second measurement (Corrier & DeLoach, 1990). Lymphoid organs, including the spleen and bursa of Fabricius, were evaluated after slaughter on day 42 of the experiment.

At 40 days old, clean stainless steel collection trays were placed under each cage, and the excreta of the birds was collected for 48 hours. A subsample of excreta was collected in polyethylene bags and stored at -20 °C to determine polyphenol content.

At 42 days old, fifteen birds per treatment were euthanized with carbon dioxide (100%). The ileum was quickly dissected and the content expressed by gentle manipulation into a plastic container and stored at -20 °C. Digesta were pooled from three birds of each replicate in the same treatment. Ileal contents were ground (1 mm screen) and used to determine the polyphenol content. On day 42, two chicks from each replicate were killed after overnight fasting to limit intestinal throughput. The whole length of the small intestine was removed, and samples were taken from the duodenum, jejunum, and ileum. The samples were flushed gently twice with physiological saline (1% NaCl) to remove intestinal contents and placed in 10% formalin for fixation. The 5-6 µ sections were made by the paraffin embedding method and stained with haematoxylin-eosin. Micrometrical analysis was carried out with a Dino-Lite digital microscope, digital Dino-Lite eye-piece and Dino-Capture 2 software on microphotographs. The variables that were measured were villus height, crypt depth and thickness of the muscularis layer. The ratio of villus height to crypt depth was calculated (Hashemi *et al.*, 2012).

Ten birds/replicate on day 42 were selected and killed by cervical dislocation. The carcasses were subsequently opened and the entire gastrointestinal tract was removed aseptically. Samples (1g) from the contents of the ceca were collected immediately and added to 5 ml glycerine in glass containers. Digesta samples were diluted in a 0.85% sterile saline solution to enumerate *Lactobacilli* and *Escherichia coli* by

conventional microbiological techniques using selective agar media as described by Jin *et al.*(1998). All microbiological analyses were conducted in duplicate, and the average values were used in the statistical analysis. *Lactobacillus* bacteria were grown on Rogosa SL agar (Merck, Darmstadt, Germany), and *Escherichia coli* was grown on McConkey agar (Merck, Darmstadt, Germany). Selective agar used to enumerate *Lactobacillus* spp. was incubated anaerobically for 48 hours at 37 °C, and selective agar used to enumerate *Escherichia coli* was incubated aerobically for 24 hours at 37 °C. Bacterial numbers were expressed as log<sub>10</sub> cfu/g of caecal digesta.

On day 42, two chicks from each replicate were killed and samples were taken from the liver of each bird. Liver weights were measured and all data are presented based on percentage of live weight of each bird. Then the liver tissue samples were collected and samples were fixed in 10% formalin and transmitted to the histological laboratory to detect histopathological changes. The 5 µ sections were made by the paraffin embedding method and stained with haematoxylin-eosin. The micrometrical analysis (diameters of cells and nucleus) was done with the Dino-Lite digital microscope, digital Dino-Lite eyepiece and Dino-Capture 2 software on microphotographs.

Data were analysed in a completely randomized design using the general linear model (GLM) procedures of SAS (SAS Institute Inc, 2001). When the differences were significant ( $P < 0.05$ ), mean values between treatments were compared using the Duncan test.

## Results

The effects of various dietary treatments on broiler BW, ADG, ADFI, and F/G are shown in Table 3. At 10 days old, the BW of birds fed the diet supplemented with GP was lower ( $P < 0.05$ ) compared with the control group. Furthermore, broilers fed the diet supplemented with GP had lower ( $P < 0.05$ ) ADG in the

**Table 3** Response of broiler chickens, in terms of bodyweight, average daily gain, average daily feed intake and feed-to-gain ratio

Item	Dietary treatments <sup>1</sup>				SEM	P-value
	Control	C+10 GP	C+ 10 GP+T1	C+10 GP+T2		
BW, g						
10 d	248.32 <sup>a</sup>	214.32 <sup>b</sup>	202.64 <sup>b</sup>	217.04 <sup>b</sup>	5.836	0.022
24 d	979.68	902.61	910.87	912.96	15.754	0.291
42 d	2416.20	2272.46	2275.92	2358.74	40.280	0.560
ADG, g						
Starter (0–10 d)	20.52 <sup>a</sup>	17.15 <sup>b</sup>	15.96 <sup>b</sup>	17.17 <sup>b</sup>	0.580	0.019
Grower (11–24 d)	52.24	49.16	50.58	49.70	1.089	0.796
Finisher (25–42 d)	79.80	76.09	75.83	80.31	2.273	0.867
Overall (0–42 d)	56.50	53.08	53.16	55.13	0.959	0.560
ADFI, g						
Starter (0–10 d)	32.80	24.60	23.37	21.88	1.587	0.052
Grower (11–24 d)	101.68	93.96	98.17	97.34	1.544	0.392
Finisher (25–42 d)	161.95	150.30	151.45	165.28	5.435	0.732
Overall (0–42 d)	113.46	101.90	104.26	109.24	2.087	0.207
Feed/gain g						
Starter (0–10 d)	1.60	1.42	1.46	1.26	0.058	0.235
Grower (11–24 d)	1.96	1.91	1.94	1.96	0.026	0.933
Finisher (25–42 d)	2.03	1.98	1.98	2.05	0.038	0.909
Overall (0–42 d)	2.00	1.92	1.96	1.98	0.015	0.241

<sup>a,b</sup>Means in the same row without a common superscript differ significantly ( $P < 0.05$ )

<sup>1</sup>Diets supplemented (+) with 10% grape pomace (10GP) and tannase enzyme (500 mg/kg of tannase (T1) and 1000 mg/kg of tannase (T2))

BW: bodyweight; ADG: average daily gain; ADFI: average daily feed intake

starter (days 0–10) period compared with the control group. In contrast, the addition of GP and tannase had no significant effect on BW, ADG, ADFI, and F/G during the grower (days 11–24) and finisher (days 25–42) experimental periods.

The main effects of the diets on carcass characteristics, abdominal fat, and yields of broilers at 42 days old are presented in Table 4. There were no significant ( $P > 0.05$ ) effects of the diets on carcasses, abdominal fat, livers or relative heart weights of broilers.

**Table 4** The relative weights of meat parts, liver, heart and abdominal fat of broiler fed different diets at 42 days of age

Dietary treatments <sup>1</sup>	Carcass	Breast	Thigh	Liver	Heart	Abdominal fat
	(% of live weight)					
Control	67.00	27.17	18.79	2.26	0.431	1.60
C+10 GP	65.89	29.31	17.84	2.15	0.432	1.59
C+10 GP+T1	65.75	26.50	18.11	2.16	0.374	1.40
C+10 GP+T2	64.06	25.79	18.50	2.26	0.432	1.62
SEM	0.598	0.588	0.178	0.052	0.015	0.093
P-value	0.405	0.169	0.258	0.801	0.478	0.853

<sup>1</sup> Diets supplemented (+) with 10% grape pomace (10GP) and tannase enzyme (500 mg/kg of tannase (T1), and 1000 mg/kg of tannase (T2))

The effects of the dietary treatments on plasma biochemical parameters are reported in Table 5. Feeding GP and tannase diets decreased serum TG ( $P < 0.01$ ). Broilers fed the diet containing 10% GP+500 mg/kg tannase had the lowest TG concentrations (60.71 mg/dl) of the experimental groups (Table 5). In comparison with the control group, a significant reduction ( $P < 0.05$ ) in LDL-C levels was observed for GP-fed groups. Broilers fed the diet containing 10% GP had lower LDL-C concentration (11.85 mg/dl). HDL-C level was increased ( $P < 0.05$ ) by dietary GP supplementation and broilers fed the diet containing 10% GP had the highest HDL-C concentration (102.93 mg/dl). There were no significant differences in serum parameters, including total protein, uric acid, glucose and cholesterol concentrations, in any of the treatments (Table 5).

**Table 5** Effects of grape pomace and tannase supplementation in chicken diets on serum biochemical parameters

Dietary treatments <sup>1</sup>	Total protein (g/dl)	Glucose (mg/dl)	Total cholesterol (mg/dl)	TG (mg/dl)	HDL-C (mg/dl)	LDL-C (mg/dl)	Uric acid (mg/dl)
Control	5.13	226.86	128.57	97.29 <sup>a</sup>	68.20 <sup>b</sup>	18.71 <sup>a</sup>	5.28
C+10 GP	5.52	219.29	107.00	63.57 <sup>b</sup>	102.93 <sup>a</sup>	11.85 <sup>b</sup>	5.20
C+10 GP+T1	5.10	214.14	109.85	60.71 <sup>b</sup>	88.55 <sup>ab</sup>	14.21 <sup>b</sup>	4.21
C+10 GP+T2	5.47	221.00	117.00	61.71 <sup>b</sup>	80.93 <sup>ab</sup>	14.71 <sup>ab</sup>	4.64
SEM	0.144	3.910	2.20	4.205	4.413	0.805	0.398
P-value	0.659	0.740	0.113	0.009	0.032	0.015	0.775

<sup>a,b</sup> Means in the same column without a common superscript differ significantly ( $P < 0.05$ )

<sup>1</sup> Diets supplemented (+) with 10% grape pomace (10GP) and tannase enzyme (500 mg/kg of tannase (T1) and 1000 mg/kg of tannase (T2))

TG: triglyceride; HDL-C: high-density lipoprotein cholesterol; LDL-C: low-density lipoprotein cholesterol

The effects of dietary treatments on SOD, GSH-Px activities and MDA concentration in serum of broilers at 42 days old are shown in Table 6. In contrast with the control group, supplementation with GP elevated ( $P < 0.05$ ) SOD and GSH-Px ( $P < 0.01$ ) activities and depressed ( $P < 0.01$ ) MDA in serum (Table 6).

**Table 6** Concentration of plasma superoxide dismutase, glutathione peroxidase and malondialdehyde in chickens fed diets supplemented with grape pomace and tannase

Items	Dietary treatments <sup>1</sup>				SEM	P-value
	Control	C+ 10 GP	C+ 10 GP+T1	C+10 GP+T2		
SOD(U/ml)	103.63 <sup>b</sup>	142.70 <sup>a</sup>	133.23 <sup>ab</sup>	164.01 <sup>a</sup>	7.072	0.014
GSH-Px(U/ml)	112.57 <sup>c</sup>	133.86 <sup>bc</sup>	147.71 <sup>b</sup>	182.31 <sup>a</sup>	7.082	0.001
MDA(nmol/ml)	7.84 <sup>a</sup>	5.34 <sup>b</sup>	4.20 <sup>c</sup>	4.13 <sup>c</sup>	0.319	0.001

<sup>a,b,c</sup> Means in the same row without a common superscript differ significantly ( $P < 0.05$ )

<sup>1</sup> Diets supplemented (+) with 10% grape pomace (10GP) and tannase enzyme (500 mg/kg of tannase (T1) and 1000 mg/kg of tannase (T2))

SOD: superoxide dismutase; GSH-Px: glutathione peroxidase; MDA: malondialdehyde

The effects of dietary treatment on antibody production, hypersensitivity and lymphoid organ weights in broilers are shown in Tables 7 and 8. Antibody titers against Newcastle disease virus at 42 days old were higher ( $P < 0.05$ ) in broilers fed the diet containing 10% GP than in those fed the control diet. The addition of GP to the diets had a significant impact on the anti-SRBC titers of total and IgM antibodies at 21 days old. In contrast, IgG antibodies were not affected by dietary treatments (Table 7).

**Table 7** Effects of grape pomace and tannase supplementation in chicken diets on antibody production (log<sub>2</sub>)

Items	Dietary treatments <sup>1</sup>				SEM	P-value
	Control	C+10 GP	C+10 GP+T1	C+10 GP+T2		
NDV antibody titer						
21 d	4	4.2	3.9	3.7	0.147	0.699
42 d	4.4 <sup>b</sup>	6.2 <sup>a</sup>	5.5 <sup>ab</sup>	5 <sup>ab</sup>	0.237	0.044
SRBC antibody titer						
Primary titer	2.40 <sup>b</sup>	3.40 <sup>a</sup>	2.60 <sup>b</sup>	2.60 <sup>b</sup>	0.142	0.048
IgG	1.40	1.60	1.60	1.40	0.114	0.879
IgM	1.00 <sup>b</sup>	1.80 <sup>a</sup>	1.00 <sup>b</sup>	1.20 <sup>b</sup>	0.099	0.002
Secondary titer	2.40 <sup>b</sup>	4.30 <sup>a</sup>	3.20 <sup>b</sup>	3.00 <sup>b</sup>	0.213	0.010
IgG	0.60 <sup>b</sup>	2.70 <sup>a</sup>	1.50 <sup>b</sup>	1.40 <sup>b</sup>	0.208	0.002
IgM	1.80	1.60	1.70	1.60	0.083	0.814

<sup>a,b</sup> Means in the same row without a common superscript differ significantly ( $P < 0.05$ )

<sup>1</sup> Diets supplemented (+) with 10% grape pomace (10GP) and tannase enzyme (500 mg/kg of tannase (T1) and 1000 mg/kg of tannase (T2)); NDV: Newcastle disease virus; SRBC: sheep red blood cell

In the secondary response, there was an increase ( $P < 0.05$ ) in total anti-SRBC titers and IgG antibodies in broilers fed the diet containing 10% GP compared with the other experimental groups. In contrast, during the secondary challenge, the SRBC-specific IgM titer in the broiler serum showed no differences across treatments. Phytohaemagglutinin injection increased ( $P < 0.05$ ) toe web thickness within 24 and 48 hours of injection in birds fed the diet containing 10% GP+500 mg/kg tannase compared with the control group (Table 8). In the present study, the relative immune organ weights were not affected by dietary treatments.

**Table 8** Effects of grape pomace and tannase supplementation in chicken diets on hypersensitivity and lymphoid organ weights

Items	Dietary treatments <sup>1</sup>				SEM	P-value
	Control	C+10 GP	C+10 GP+T1	C+10 GP+T2		
Hypersensitivity (mm), d 38						
24 hours after	0.87 <sup>b</sup>	0.81 <sup>b</sup>	1.23 <sup>a</sup>	0.77 <sup>b</sup>	0.066	0.045
48 hours after	0.63 <sup>b</sup>	0.64 <sup>b</sup>	0.94 <sup>a</sup>	0.60 <sup>b</sup>	0.048	0.045
Lymphoid organ weight (g/kg BW)						
Spleen	0.106	0.107	0.092	0.103	0.005	0.731
Bursa of Fabricius	0.048	0.045	0.062	0.050	0.003	0.463

<sup>a,b</sup>Means in the same row without a common superscript differ significantly ( $P < 0.05$ )

<sup>1</sup>Diets supplemented (+) with 10% of grape pomace (10GP) and tannase enzyme (500 mg/kg of tannase (T1) and 1000 mg/kg of tannase (T2))

The total ileal and excreta polyphenol contents of birds fed diets containing GP and tannase are reported in Table 9. Birds fed GP diets showed higher ileal and excreta polyphenol contents than those fed control diets. The inclusion of tannase in GP diets increased ileal and excreta polyphenol content, being higher in animals fed diets containing 10% GP+ 500 mg/kg tannase.

**Table 9** Effect of diets containing grape pomace and tannase on ileal and excreta total polyphenol content expressed as mg gallic acid equivalent per gram of dry matter

Dietary treatments <sup>1</sup>	Total polyphenol (mg GAE/g DM)	
	Ileal	Excreta
Control	2.139 <sup>b</sup>	5.179 <sup>c</sup>
C+10 GP	5.867 <sup>a</sup>	6.379 <sup>b</sup>
C+10 GP+T1	6.044 <sup>a</sup>	7.991 <sup>a</sup>
C+10 GP+T2	5.954 <sup>a</sup>	7.429 <sup>a</sup>
SEM	0.468	0.276
P-value	0.011	0.001

<sup>a,b,c</sup>Means in the same column without a common superscript differ significantly ( $P < 0.05$ )

<sup>1</sup>Diets supplemented (+) with 10% of grape pomace (10GP) and tannase enzyme (500 mg/kg of tannase (T1) and 1000 mg/kg of tannase (T2))

GAE: gallic acid equivalent

The histomorphometrical measurement results of the various parts of small intestine of broiler chicks are shown in Table 10. The muscularis thickness of the chicken duodenum was decreased in the Trial 3 compared with the control groups ( $P < 0.05$ ), but villus height and crypt depth were not affected. In the jejunum parts of small intestine, villus height and villus height/crypt depth ratio were reduced ( $P < 0.01$ ) in birds fed the GP diet compared with control groups (Table 10). There were no significant effects of diets on villus height and the villus height/crypt depth ratio in the ileum ( $P > 0.05$ ). Muscularis thickness and crypt depth were decreased in birds fed GP supplemented with tannase compared with those fed the control.

**Table 10** Effects of grape pomace and tannase supplementation in chicken diets on morphology of the small intestine at 42 days of age

Items	Dietary treatments <sup>1</sup>				SEM	P-value
	Control	C+10 GP	C+10 GP+T1	C+10 GP+T2		
<i>Duodenum</i>						
Villus height (µm)	1295.9	927.8	1071.3	1263.3	68.290	0.176
Crypt depth (µm)	296.37	299.20	249.45	310.06	11.198	0.239
Villus height/crypt depth	4.36	3.15	4.33	4.09	0.223	0.183
Muscularis thickness (µm)	149.86 <sup>a</sup>	110.25 <sup>ab</sup>	84.28 <sup>b</sup>	98.11 <sup>b</sup>	9.274	0.038
<i>Jejunum</i>						
Villus height (µm)	1198.5 <sup>a</sup>	700.6 <sup>b</sup>	682.41 <sup>b</sup>	647.43 <sup>b</sup>	74.357	0.001
Crypt depth (µm)	300.29	262.88	283.94	266.27	15.079	0.850
Villus height/crypt depth	3.98 <sup>a</sup>	2.68 <sup>b</sup>	2.46 <sup>b</sup>	2.43 <sup>b</sup>	0.201	0.001
Muscularis thickness (µm)	143.81	132.63	94.68	134.27	10.496	0.406
<i>Ileum</i>						
Villus height (µm)	665.19	627.07	517.81	709.41	32.018	0.171
Crypt depth (µm)	285.12 <sup>a</sup>	211.31 <sup>b</sup>	193.81 <sup>b</sup>	248.63 <sup>ab</sup>	13.453	0.041
Villus height/crypt depth	2.40	2.98	2.68	2.84	0.136	0.540
Muscularis thickness (µm)	147.14 <sup>a</sup>	95.83 <sup>b</sup>	96.65 <sup>b</sup>	94.14 <sup>b</sup>	7.103	0.002

<sup>a,b</sup>Means in the same row without a common superscript differ significantly ( $P < 0.05$ )

<sup>1</sup>Diets supplemented (+) with 10% of grape pomace (10GP) and tannase enzyme 500 mg/kg of tannase (T1) and 1000 mg/kg of tannase (T2)

Bacteriological analysis results at 42 days old are detailed in Table 11. In the caecal digesta, the inclusion of GP in the chicken diets reduced ( $P < 0.05$ ) the concentration of *Escherichia coli* and increased ( $P < 0.05$ ) that of *Lactobacillus* compared with the control diet (Table 11). The caecal populations of *Lactobacillus* were higher in broilers fed the 10 % GP or 10% GP + 500 mg/kg tannase diet compared with birds fed the control diet at 42 days old. In contrast, Trial 4 broilers exhibited lower caecal *Escherichia coli* counts than other experimental groups.

**Table 11** Effects of grape pomace and tannase supplementation in chicken diets on caecal bacterial content at 42 days of age

Microbial activity (cfu) <sup>2</sup>	Dietary treatments <sup>1</sup>				SEM	P-value
	Control	C+ 10 GP	C+ 10 GP+T1	C+10 GP+T2		
Lactobacillus	6.53 <sup>b</sup>	7.12 <sup>a</sup>	7.03 <sup>ab</sup>	6.54 <sup>b</sup>	0.100	0.041
Escherichia coli	6.57 <sup>a</sup>	6.51 <sup>a</sup>	6.73 <sup>a</sup>	5.96 <sup>b</sup>	0.106	0.045

<sup>a,b</sup>Means in the same row without a common superscript differ significantly ( $P < 0.05$ )

<sup>1</sup> Diets supplemented (+) with 10% of grape pomace (10GP) and tannase enzyme (500 mg/kg of tannase (T1) and 1000 mg/kg of tannase (T2))

<sup>2</sup>CFU:colony-forming units are expressed as log10 colonies per gram of caecal contents

The effects of dietary treatment on liver histology and concentrations of liver function indicators AST and ALT are summarized in Table 12. A significant statistical elevation ( $P < 0.05$ ) in the activity of AST and ALT enzymes was observed in GP-fed groups compared with the control.

**Table 12** Effects of grape pomace and tannase supplementation in chicken diets on liver histomorphometric parameters, aspartate aminotransferase and alanine aminotransferase in plasma

Item	Dietary treatments <sup>1</sup>				SEM	P-value
	Control	C+ 10 GP	C+ 10 GP+T1	C+10 GP+T2		
Diameters of cells ( $\mu\text{m}$ )	8.24	7.98	8.31	8.31	0.119	0.784
Diameters of nucleus ( $\mu\text{m}$ )	4.43	4.81	4.63	4.65	0.088	0.563
ALT(u/l)	12.42 <sup>b</sup>	11.57 <sup>b</sup>	13.28 <sup>ab</sup>	15.85 <sup>a</sup>	0.586	0.047
AST(u/l)	193.57 <sup>b</sup>	333.00 <sup>a</sup>	352.00 <sup>a</sup>	316.43 <sup>a</sup>	19.196	0.007

<sup>a,b</sup>Means in the same row without a common superscript differ significantly ( $P < 0.05$ ).

<sup>1</sup> Diets supplemented (+) with 10% of grape pomace (10GP) and tannase enzyme (500 mg/kg of tannase (T1) and 1000 mg/kg of tannase (T2)

ALT:alanine aminotransferase; AST: aspartate aminotransferase.

No significant ( $P > 0.05$ ) histopathological changes were seen for birds fed GP and different levels of tannase compared with the control group. There were no significant ( $P > 0.05$ ) effects of feeding diets on liver weights in broilers (Table 4).

## Discussion

There are few references in the literature to the use of grape by-products in chicken feed. Hughes *et al.* (2005) and Lau & King (2003) reported growth depression in chickens fed diets containing grape seed extract (GSE). In contrast, Hajati *et al.* (2015) reported that GSE supplementation at the levels of 300 mg/kg diet improved the bodyweight of broilers. Recent works by Brenes *et al.* (2008) and Chamorro *et al.* (2013) showed that concentrations of up to 6% GP and 0.25% GSE could be used in chicken feed without modification in performance, in the size of the digestive organs or the ileal digestibility of protein and in amino acids. Grape pomace contains high levels of fibre and polymeric polyphenols as procyanidins with the capacity to bind and precipitate dietary and endogenous proteins. Therefore, the incorporation of GP at high doses in chicken diets might impair nutrient digestion and growth. The utilization of enzymes with the capacity to hydrolyse complex polyphenols in GP might allow the use of higher doses of GP in chicken diets. Thus, in the present study, high doses of GP were used. Because all diets were formulated to contain the same fibre content, differences should be attributed to the polyphenol content.

The results of the current study showed that dietary inclusion of up to 10 percent GP did not affect chick growth performance. Consequently, the use of the tannase enzyme did not enhance the parameters. In agreement with the current results, Chamorro *et al.* (2015) reported that the inclusion of enzymes (carbohydrases and tannase, 500 mg/kg) in chicken diets containing GP (5% and 10%) modified the polyphenol polymeric structures in the ileal content, with increased concentrations of gallic acid, catechin and epicatechin in birds fed carbohydrases, and of gallic acid, catechin, epicatechin, procyanidins PB1 and PB2 and epicatechin gallate. These findings show that the inclusion of enzymes in diets containing GP increased the amount of total polyphenol released in the intestine, although this effect was not accompanied by an increase in performance. There were no significant effects of the diets on carcasses, abdominal fat and relative liver or heart weights in broilers. These results are in agreement with those of Brenes *et al.* (2008), who reported that the inclusion of graded concentrations of GP did not affect relative abdominal fat, liver, pancreas, and spleen weight.

In this study, serum total protein and cholesterol levels were unchanged. This agrees with studies that reported that including plant extracts rich in polyphenols such as rosemary, grape, citrus and marigold at 5 g/kg diet in rats (Gladine *et al.*, 2007) and inclusion of GP in laying hen diets up to 6 percent by Kara *et al.* (2015) has no effect on serum protein and cholesterol. Hajati *et al.* (2015) reported that GSE supplementation (150, 300, 450 mg/kg) decreased the concentration of serum glucose. In contrast, in the current study, GP supplementation of chicken diets did not have a significant impact on serum glucose. In agreement with the current results, Ozgan (2008) stated that the plasma glucose level was not affected by the inclusion of 1.0% grape seed oil in laying hen diets. Similar to the results of this study, administration of GSE reduced blood levels of total TG, LDL-C and increased HDL-C in broilers (Hajati *et al.*, 2015). The concentration of uric acid in broiler serum was not affected by dietary treatments ( $P > 0.05$ ). Antioxidant activity is the most notable bioactivity of phenolic compounds from GP (Xia *et al.*, 2010; Georgiev *et al.*,

2014). *In vitro* and *in vivo* studies have shown that flavonoids in wine by-products behave like free radical scavengers by acting as powerful antioxidants and metal chelators. This inhibits formation of the superoxide ion and indirectly inhibits redox-sensitive transcription factors and pro oxidant enzymes (Puiggros *et al.*, 2005). Polyphenols also activate antioxidant enzymes, reduce a tocopherol radical (tocopheroxyls), inhibit oxidases, and increase levels of uric acid and substances of low molecular weight. The antioxidative properties of polyphenols result mainly from their ability to donate hydrogen from hydroxyl groups positioned along the aromatic ring in order to terminate the free radical oxidation of lipids and other biomolecules (Foti *et al.*, 1994).

The current findings are consistent with those of Seven *et al.* (2008), who reported that feeding plant extracts rich in phenolic compounds resulted in an increase in blood GSH-Px activity of chronic heat-exposed broiler chickens. Similar positive changes in plasma MDA and SOD levels to those in the current study were reported in a study of layer chickens fed with citrus flavonoids (Lien *et al.*, 2008). Results from this study confirm that dietary GP can delay lipid oxidation in plasma and reduce the potential risk induced by lipid oxidation. In the present study, the addition of tannase to diets containing GP reversed the beneficial effect observed in GP diets. Birds fed with GP diet supplemented with 1000 mg/kg tannase showed higher SOD and GSH-Px activities and lower MDA concentration in comparison with other experimental groups (Table 6).

The addition of GP and tannase in the chicken diets could increase the cellular and humoral immune responses. In poultry production, it is important to improve immunity to prevent infectious diseases. A variety of factors such as vaccination failure, infection by immune-suppressive diseases, and abuse of antibiotics can induce immunodeficiency. Grape pomace is particularly rich in a wide range of polyphenols. Konowalchuk & Speirs (2009) reported that grape seed proanthocyanidins show up to 50 times more antioxidant activity than vitamin E. Immunity is influenced by oxidative stress and improving the antioxidant status of the animal may enhance its immunity. Similar to the current data, Iqbal *et al.* (2015) found that broilers fed the grape polyphenols diets showed higher ( $P < 0.05$ ) antibody titers against ND virus than those fed the control diet. Additionally, Acamovic & Brooker (2005) reported immunostimulating activity of polyphenols and oregano essential oil with respect to the mononuclear phagocyte system, cell, and humoral immunity. Today, there is compelling evidence that suggests that the mechanisms by which plant polyphenols exert their properties, such a mode of action would induce the inhibition of key enzymes, modulate cell receptors or transcription factors, and perturb protein (or peptide) aggregates, which can regulate cell functions related for example to growth and proliferation, inflammation, apoptosis, angiogenesis, metastasis, and immune responses by affecting signal transduction pathways (Zhu *et al.*, 1997; Yang *et al.*, 2009).

Immune organs, including the spleen and bursa of Fabricius, are considered important sites for antigen recognition and activation of immune cells (Akter *et al.*, 2006). The relative weight of the spleen and bursa reflect the host's ability to produce T and B lymphocytes during an immune response. In the present study, the relative immune organ weights were not significantly affected by dietary treatments. In agreement, Brenes *et al.* (2008) and Rahimi *et al.* (2011) showed no significant differences in the relative weight of the spleen and bursa in broilers fed a diet containing GP concentrate and thyme compared with the control groups. The biological effect of polyphenols depends on their availability. Monomeric and some oligomeric polyphenols have been found to be absorbed in the small intestine, while polymeric forms are poorly absorbed, and must be hydrolysed by the intestinal microbiota (Gonthier *et al.*, 2003). As expected, birds fed GP diets showed higher intestinal (ileal and excreta) total polyphenol contents than those fed control diets. Recent studies have estimated that the amount of non-absorbable polyphenols reaching the colon is high and microbe-derived phenolic metabolites excreted in urine represent the largest proportion of polyphenol intake (Monagas *et al.*, 2010).

Birds lack a specific urinary excretory system. Consequently the excreta of chicks are composed of urine and undigested compounds. Thus, the total polyphenols that were determined in excreta in this experiment included phenolic substances from undigested and microbe-derived phenolic metabolites. This might explain why the amount of phenolic compounds in excreta was higher than those obtained from the ileal content. The authors hypothesized that the use of enzyme with tannase activities might hydrolyse the complex polyphenol structure of grape pomace into more simple phenols, enhancing the amount of bioactive available substances and, therefore, its antioxidant capacity. Lafay & Gil-Izquierdo (2008) reported that low molecular weight phenolic compounds can be much more easily absorbed and might contribute more directly to beneficial health effects.

Mucosa status and their microscopic structure can be good indicators of the response of the intestinal tract to active substances in feeds. Few reports have documented the effects of dietary polyphenols or related phenolics on localized intestinal growth and function in broiler chickens and their contribution to changes in performance. In a study in which chicks and rats were given diets high in freeze-dried tannins

extracted from fava beans (*Vicia faba*), Ortiz *et al.* (1994) found histological lesions in the ileum, suggesting a loss in absorptive capacity. Results obtained by Sell *et al.* (1985), using high and low tannin sorghum in rats and laying hens, revealed milder negative effects. Examination of the distal digestive tract under a light microscope showed normal cells.

The only effect appeared to be a mild reduction in crypt depth and intestinal wall thickness and an increase in mucus secretion in the intestinal tract. In the present study, villus height and crypt depth were reduced, particularly in the jejunum. It is assumed that increase of the villus height is paralleled with increasing digestive and absorptive function of the intestine as a result of increased absorptive surface area, expression of brush border enzymes, nutrient transport systems (Caspary, 1992), and an increased bodyweight gain (Zijlstra *et al.*, 1996). Sehm *et al.* (2007) reported that the inclusion of red wine pomace had an inhibitory effect on the jejunum villi growth. In agreement with the current results, Laurent *et al.* (2005) observed a decrease in the size of the microvilli from the addition of GSE in human intestinal Caco-2 cells. Viveros *et al.* (2011) reported that inclusion of grape pomace concentrate (GPC) and GSE decrease villus height and crypt depth in broiler chickens. Intestinal cell proliferation occurs mainly in the crypts (Geyra & Sklan, 2001). Thus, the large crypt suggests a high nutrient requirement for intestinal maintenance and reduced efficiency of the bird. In this view, the lower crypt depth as a result of fed GP can be considered as a beneficial way to decrease the cost of intestinal maintenance in birds.

In the present study, a decrease in villus height/crypt depth ratio in birds fed diets was also found. A lengthening of the villus and a short crypt can lead to better nutrient absorption, decreased secretion in the gastrointestinal tract, increased disease resistance, and greater overall performance. In this sense, the low villi height: crypt depth ratio could have caused poorer growth in GP-fed chicks. In contrast, an increase in this ratio in the jejunum in birds fed GPC diets was reported by Viveros *et al.* (2011). Moreover, muscularis thickness was decreased in birds fed a GP diet. In contrast, Viveros *et al.* (2011) reported that muscularis thickness was increased in birds fed GPC and GSE diets. Inclusion of feed antibiotics in diets of chicken reportedly leads to decreased thickness of walls and reduced weight (Miles *et al.*, 2006). Many *in vitro* studies have demonstrated the antimicrobial properties of polyphenols (Arunachalam *et al.*, 2003). Improved intestinal morphology characteristics such as lower crypt depth and epithelial thickness could lead to increased feed utilization and improved performance. There have been few studies on the interaction of polyphenol compounds with intestinal microbiota in animal nutrition. Numerous *in vitro* studies (Papadopoulou *et al.*, 2005; Ganan *et al.*, 2009) have shown that flavonoids in grape by-products have the capacity to inhibit the growth of organisms such as *Staphylococcus aureus*, *Escherichia coli*, *Candida albicans* and *Campylobacter*. Likewise *in vivo* studies have shown that resveratrol has potential as an antibiotic alternative for reversing the adverse effects of weaning stress on the growth performance, immunity and microbial environment in *E. coli* and *Salmonella*-challenged piglets (Ahmed *et al.*, 2013).

The inhibiting effect of polyphenolic compounds on bacteria could be because of mechanisms related to their capacity to adhere to cellular membranes, to interact with bacterial enzymes and to sequester metallic ions from the substrate (Scalbert, 1991; Cushnie & Lamb, 2011). In the present experiment, the *lactobacilli* stimulant activity of GP in the cecum could be attributed to the composition and phenolic profile of these by-products and could depend directly on extractable polyphenol contents. A possible explanation for the stimulatory effect of polyphenolic compounds on bacterial growth is that some microorganisms are able to use these compounds as nutritional substrates. In *lactobacilli*, these bacteria possess the ability to metabolize phenolic compounds supplying energy to cells and positively affecting the bacterial metabolism (Garcia-Ruiz *et al.*, 2008). In agreement with the current results, Viveros *et al.* (2011) confirmed the antibacterial effect of polyphenols in GPC and GSE on certain intestinal bacteria. This effect differs depending on the segment analysed (ileum or caecum). The inclusion of these by-products in bird diets exerted an antimicrobial effect on *Clostridium* in the ileum, while in the caecum it was associated with an increase in populations such as *Lactobacillus* and *Enterococcus*.

Regarding histopathological examination, Iqbal *et al.* (2015) reported that no changes were observed in the livers and kidneys of birds fed low and medium levels of GP polyphenols in diets. Also, mild changes in the livers and kidneys of birds fed a diet with a high level of GP polyphenols suggested that even higher polyphenols levels were relatively nontoxic to birds (Iqbal *et al.*, 2015). However, Haseeb *et al.* (2011) reported protective effects of GSE against aluminium toxicity in rats. Similarly, Sun *et al.* (1999) reported alleviating effects of GP polyphenols on chronic ethanol-induced morphological changes in rats. The GSE was reported to have many other beneficial effects such as hepatoprotection (Hassan, 2012), and protection against silica-induced pulmonary fibrosis (Ali *et al.*, 2008), and ethanol-induced oxidative damage and ulcer (Cuevas *et al.*, 2011). In the present study, GP supplementation to chicken diets did not have a significant impact on biochemical parameters. Only AST and ALT activity were increased significantly in tannase supplementation groups compared with the control ( $P < 0.05$ ).

Injury to the liver results in the release of intracellular enzymes into the extracellular space (Kristensen, 1994). Increases in plasma levels of the liver function enzymes AST and ALT are well known diagnostic indicators of liver injury (Chiu *et al.*, 2013). The activity of AST is currently considered a sensitive indicator of hepatocellular disease in avian species. The greater plasma AST concentration in the Trial 3 and ALT in the Trial 4 relative to the other treatment may suggest hepatocellular necrosis by polyphenol compounds, which causes an increase in permeability of the cell membrane, resulting in the release of transaminase in the bloodstream. In agreement with the current results, Lu *et al.* (2014) reported that plasma AST concentration in a high-oxidant diet with vitamin E level at NRC requirement of 10 IU/kg was greater on day 21 and 42 relative to the group that were fed standard control diet supplementation with vitamin E level at NRC requirement of 10 IU/kg. Khudiar & Aldabaj (2015) reported that grape seed oil (GSO) supplementation could alleviate hepatic changes induced in adult male rabbits by sodium fluoride as reflected by normalization of the measured liver function markers suggesting its potential hepatoprotective effect. The natural antioxidant in GSO may be responsible for its antioxidant property and for protection of DNA damage (Hassan *et al.*, 2014) and was claimed to be the mechanism of hepatoprotective activity of GSO. Degenerative changes in liver were reported on feeding high tannin sorghum to poultry. Sannamani (2002) reported significant pathological changes in most of the tissues (lungs, liver, kidney, heart, intestine and pancreas) of birds fed on a 100% red sorghum-based diet without supplemental methionine. Moreover, histopathological examination of the liver of chicks or rats fed on the diets containing tannin showed hydropic degeneration of hepatocytes (Ortiz *et al.*, 1994). There was no significant effect of feeding diets on relative liver weights in broilers. These results are in agreement with the those of Brenes *et al.* (2008), who reported that the inclusion of graded concentrations of GPC did not affect the relative liver and spleen weight.

Many investigators have demonstrated the efficacy of grape seed proanthocyanidins and grape juice as an inhibitor of lipid peroxidation and as a powerful free radical scavenger *in vitro* and *in vivo* (Dani *et al.*, 2008; Khalifa *et al.*, 2011; Naguib, 2011) in addition to its hepatoprotective effect (Hassan & Al-Rawi, 2013). Thus it was suggested that the active component of grapes protect the hepatocytes from liver damage and subsequent leakage of enzymes into the circulation, and may have a curative effect against elevated transaminases (Ahmed & Fatani, 2007; Al-Attar, 2015). Thus, like many plant extracts, grape polyphenols have been shown to play a double-edged role in regulating the inflammatory system of the host. Caution in dosage should be taken carefully to avoid switching from beneficial effects to adverse ones. Blood proteins are important complementary constituents in the diagnosis of hepatic disease, although determination of these proteins seldom leads to diagnosis, although it helps to evaluate the nature, severity and progress of disease (Lumeij, 2008). Besides, GSE causes inhibition of gluconeogenesis and prevents catabolism of protein and its conversion to glucose (Bujanda *et al.*, 2006). This may lead to increased levels of total polyphenol concentration in serum. Previous works found that GSE prevents oxidative injury (Puiggros *et al.*, 2005), modulates glucose haemostatic (Montagut *et al.*, 2010), and has an insulin-mimetic effect on adiposities and adipose tissue (Pinent *et al.*, 2004). GSE, a well-known dietary supplement, (Weber *et al.*, 2007), with its proanthocyanidins, possesses insulinomimetic properties (Decorde *et al.*, 2009) by stimulating glucose uptake in insulin-sensitive cell lines, and decreases hyperglycaemia in streptozotocin diabetic rats (Pinent *et al.*, 2004). Polyphenols are known to inhibit gut glucose absorption and peripheral tissue glucose uptake by glucose transporter (Pandy & Rizvi, 2009), protect pancreatic islets cells, and inhibit insulin resistance (Zunino *et al.*, 2007; Meydani & Hasn, 2010), while flavonoids possess antidiabetogenic activity attributed to increased pancreatic secretion and insulin release from beta cells (Sharma *et al.*, 2008; Sridhar *et al.*, 2005).

## Conclusion

The results presented in this study showed that increasing the concentration of GP up to 10 percent did not impair the performance and carcass characteristics of broiler chickens. The addition of GP to the chicken diets could increase cellular and humoral immune responses. The inclusion of tannase in diets containing GP increased the bioavailability and amount of total polyphenol released in the intestine and improved the antioxidant status of broilers. Based on the results of the present study, it can be stated that dietary polyphenol-rich grape products were effective in increasing the caecal populations of beneficial bacteria in the cecum. Crypt depth and epithelial thickness of chicken duodenum and the villus height and crypt depth were significantly reduced in the jejunum. These facts may have an important influence on the physiology and biochemistry of the gut. To optimize health benefits and minimize possible negative effects, more studies are needed to establish appropriate dosages of grape polyphenols and their bioavailability.

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### Authors' Contributions

SKE and BN were in charge of the experimental design, implementation and writing the manuscript. All authors participated in interpreting and reviewing the results of the study.

### Conflict of Interest Declaration

There are no conflicts of interest.

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