



Effects of spent mushroom compost meal on growth performance and meat characteristics of grower geese

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ABSTRACT - The objective of this study was to determine the effect of spent mushroom compost (SMC) meal on the growth performance and meat characteristics of geese. The SMC extracts contained 2.49 ± 0.62 mg gallic acid equivalent/g dry weight (DW) and 1.08 ± 0.15 mg quercetin equivalent/g DW of the total phenolic and flavonoid contents. A total of 120 White Roman geese, aged five weeks, were randomly distributed among 12 pens and fed a grower diet *ad libitum* during the growing period, with each pen containing five males and five females in a completely randomized design. Each treatment comprised three pens (total of 30 geese), including control (corn-soybean meal); supplementation with 5% SMC meal (5% SMC); supplementation with 10% SMC (10% SMC); and supplementation with 15% SMC (15% SMC), for eight weeks. The results revealed that the body weight of the 15% SMC group was significantly lower than that of the control group at the age of 12 weeks. There were no significant effects among the groups on feed conversion ratio or intake. The malondialdehyde content of the serum in the 15% SMC group was lower than that of the control group at week 12. There were no significant effects among the groups for blood biochemical parameters in grower geese at week 12. The color values of meats in the SMC groups were higher than those obtained with control group. The flavor and acceptability score of meats in the 5% SMC group were significantly higher than for the 15% SMC and control group. Supplementation with SMC at 5% in the diet has no adverse effects on the growth performance of grower geese. However, SMC meal at 5% in the diet favorably affects sensory attributes (meat flavor and acceptability).

Key Words: antioxidant, meat sensory evaluation, white Roman goose

Introduction

The cost of raw materials for poultry feeding has significantly increased annually. Thus, finding alternative feeds has become an important task of the feed/food industry. The mushroom industry is experiencing a global development, and the spent mushroom compost (SMC) from mushroom growing is typically recycled agricultural waste products that could thus provide a low-cost feed to animals.

Currently, mushrooms are typically cultivated within a 3-D architectural structure inside air-conditioned facilities. Substrates commonly used in mushroom production include agricultural by-products, such as cereal straw (rice, wheat,

corn, and barley), cotton, cobs, husks, and pulp (Chang and Miles, 1989), which undergo a process of mixing, high-temperature composting, pasteurization, inoculation with a pure mushroom culture, and incubation, prior to mushroom production. When the mushroom fruiting bodies mature and are ready for sale, the spent mushroom compost (SMC) is available as a by-product/waste (Herrero-Hernández et al., 2011; Tajbakhsh et al., 2008). At this stage, the compost is known as “spent mushroom compost” (SMC), and it also has nutritional components (Mullen and McMahon, 2001): approximately 73.6% neutral detergent fiber, 55.0% acid detergent fiber, 8.1% crude protein, 2.1% ether extract, 9.8% non-fibrous carbohydrate, and 6.4% crude ash (Kim et al., 2007). Based on this composition, SMC has potential as a feed resource for livestock. Growing Hanwoo steers were supplemented with 50% of the microbially-fermented spent mushroom substrates of the *ad libitum* group, which resulted in a tendency of increased live weight gain from 8 to 12%, as compared with the control group (Kim et al., 2012). Suwandystuti and Bata (2012) indicated that spent mushroom rice straw compost could replace 75% fresh

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Napoer grass (DM) for male cattle. Kim et al. (2012) reported that about 660,000 t of SMC (*Pleurotus eryngii*) were produced annually in Korea, while in Taiwan, at present, an increase of 130% means the annual production is about 80,000 metric tons of SMC.

In addition to nutrition composition, SMC is composed of fungal mycelia, extracellular enzymes (such as cellulase and xylanase) secreted from mushrooms for degradation of substrates, and unused lignocellulosic substrates (Kumaran et al., 1997; Ayala et al., 2011). Moreover, one of these, laccase, a kind of enzyme for a lignin-degrading enzyme system, was extracted from the spent mushroom compost in *Pleurotus eryngii* (Lim et al., 2013). Park et al. (2012a) showed that goats fed 15% SMC for six weeks had significantly higher white blood cell and lymphocyte counts than the control group or goats fed 20% SMC. Park et al. (2012b) reported that elk fed 15% or 20% SMC had higher blood urea nitrogen and glucose concentrations than the control group.

Geese are herbivorous poultry that resist coarse foods and can grow well if fed good forage grass. Hollister et al. (1984) showed that when goslings were fed 20% Kentucky blue grass and dehydrated alfalfa, the feed conversion was not significantly different from those of goslings fed a control (corn-soybean meal) diet; however, feed costs were reduced. To our knowledge, little information is presently available regarding the influence of SMC meal applied as poultry feed. Therefore, the objective of this study was to investigate the use of SMC meal as feed on the growth, antioxidant enzymes activity, and meat characteristics of grower geese.

Material and Methods

The spent mushroom compost (*Pleurotus ostreatus* (Jacq.exFr.) Kumm.) used in this study was provided by a mushroom producer in Nantou county, Taiwan. The fresh SMC was dried in a forced-air dryer at 50 °C for 24 h and then ground to a powder (approximately 2.5 mm in size) prior to its addition to the feed. The SMC extracts were added to 100% distilled water (1:10, w/v) at 95 °C for 2 h after filtering (Advantec NO. 1, Tokyo, Japan). The filtrate was evaporated until dry under vacuum conditions. The lyophilized extracts were rehydrated and the concentration was adjusted to 1 mg/mL for subsequent analysis.

Total phenolic contents were determined using a Folin-Ciocalteu reagent according to the method described by Kujala et al. (2000). The Folin-Ciocalteu reagent was mixed evenly with SMC extracts before adding the Na₂CO₃ solution, and measured with a spectrophotometer at OD₇₃₀.

Then, with the contents of the phenolic compounds of the extracts, one microgram of the gallic acid equivalent (GAE) was determined using an equation that was obtained from the standard gallic acid graph. The flavonoid content of SMC extracts was determined by following the colorimetric method (Chang et al., 2002). Briefly, 0.5 mL of SMC extracts in methanol were mixed with methanol, 10% aluminum chloride, and 1 M potassium acetate, and left for 25 min at room temperature. The absorbance of the reaction mixture was measured at OD₄₁₅ with a spectrophotometer, and the calibration curve was obtained by preparing quercetin solutions.

One hundred twenty White Roman geese aged five weeks were randomly distributed among 12 pens and fed a grower diet *ad libitum* during the growing period, with each pen containing five males and five females, arranged in a randomized complete design. Initially, each treatment comprised three pens, with 10 geese in each pen, including: control (corn-soybean meal), supplementation with 5% SMC meal (5% SMC); supplementation with 10% SMC (10% SMC); and supplementation with 15% SMC (15% SMC), for 8 weeks (from weeks 5 to 12). The care and use of all geese complied with the Regulations of Laboratory Animals and were approved by the Institutional Animal Use and Care Committee (IACUC) according to the Regulations of Laboratory Animals, Changhua Animal Propagation Station, Livestock Research Institute (CAPS-LRI, located at 23°51'N and 120°33'E), COA-LRI, Council of Agriculture, Taiwan. During the experimental period, the geese experienced a natural photoperiod and day length of 11.00 to 12.00 h. The controlled-environment finishing house was 1.26 m² per goose. The pen of the finishing house measured 3.95 × 2.5 m (9.88 m²). It contained a wire floor, one tank, and two water dispensers, and drinking water was provided *ad libitum*. The SMC contained 4.71% CP and 2,627 kcal ME/kg, respectively. During the experimental period, the geese were fed a grower diet (Table 1), which was formulated to meet the nutrient requirements according to the NRC (1994), and the proximate composition was analyzed according to AOAC (1984).

At the end of 12 weeks, the performance of the geese was assessed by measuring the feed intake and body weight (BW); BW gain and feed conversion ratio were recorded. On the same day, 12 geese (six males and six females) from each group (four geese per pen) were randomly selected, blood samples were obtained from each goose by jugular venipuncture, and then they were killed by exsanguination. The blood samples were centrifuged at 2500 × g for 30 min at 4 °C; the samples were then stored at -20 °C until analysis. The samples for

the carcass characteristics were individually weighed and harvested, and then expressed in grams per goose. The left breast muscles were frozen at -20°C for further meat characteristic evaluation.

First, the left breasts were weighed and cooked to an internal temperature of 80°C in a digital thermostat water bath until the temperature of the meat center reached 71°C , after which the breasts were cooled for 30 min (AMSA, 1995). Each of the breasts was cut into six adjacent 1 cm (width) \times 1 cm (thickness) \times 3 cm (length) strips, parallel to the direction of the muscle fibers (Gentry et al., 2002). Each strip was sheared once, and the mean was calculated for each breast. Samples were sheared perpendicular to the muscle fibers using a Stable Micro Systems TA-XT-Plus Texture Analyzer (Stable Micro Systems, UK). Cooking loss was reported as a percentage and calculated as (initial weight – final weight)/(initial weight) \times 100%. For sensory evaluation, the breast meat testing was performed by sensory panelists in triplicate on each sample. A trained ten-member panel consisting of researchers from the Division of Animal Products Processing, Livestock Research Institute, Council of Agriculture in Taiwan, was used to evaluate

the chicken patties. Panelists were trained according to a sensory evaluation procedure. Each breast was evaluated in terms of color, flavor, juiciness, tenderness, and overall acceptability. Breast samples were placed in a vacuum bag, and heated at 80°C in a water bath for 25 min, then cut into quarters (size is $5 \times 5 \times 3$ cm), and randomly served to the panelists. Each sample was coded with a randomly selected three-digit number. Sensory evaluations were performed under fluorescent lighting. Panelists were instructed to cleanse their palates between samples using warm water (30°C). The flavor (1 = extremely undesirable, 6 = slightly desirable, 10 = extremely desirable), juiciness (1 = extremely dry, 6 = slightly juicy, 10 = extremely juicy), visual color (1 = extremely undesirable, 6 = slightly desirable, 10 = extremely desirable), and overall acceptability (1 = extremely undesirable, 6 = slightly desirable, 10 = extremely desirable) of the cooked samples were evaluated using a 10-point descriptive scale. This analysis was conducted using the hedonic test, as described by Choi et al. (2010a).

The serum biochemical values of geese were measured using an automatic biochemical analyzer (Hitachi, 7150 auto-analyzer, Hitachi, Tokyo, Japan). A spectrophotometer was used to colorimetrically assay the activities of superoxide dismutase (SOD) and Trolox equivalent antioxidant capacity (TEAC) (Wheeler et al., 1990). The procedures were conducted with assay kits purchased from Cayman Chemical Co. (Ann Arbor, MI, USA). The malondialdehyde (MDA) level was detected with 2-thiobarbituric acid, and the change of absorbance at OD_{523} was monitored with a spectrophotometer. All samples were measured in triplicate, and at appropriate dilutions, to allow the activities of enzymes to achieve the linear range of standard curves. Antioxidative enzyme activities were expressed as unit (U) per milliliter of serum (Lee et al., 2012b).

The data of the variables collected were analyzed statistically using the general liner models procedure (GLM) of SAS software (Statistical Analysis System, version 9.0.1) following a random arrangement.

The mathematical model was:

$$Y_{ij} = \mu + T_i + \varepsilon_{ij}$$

where Y_{ij} = observed response of bird in a pen; μ = overall mean; T_i = fixed effect of SMC supplementation; and ε_{ij} = residual error when pen t was regarded as experimental unit t , $\varepsilon_{ij} \sim N(0, \sigma^2)$. The mean values were compared between four SMC supplementations using the LSMEANS with the significance level of $P < 0.05$.

Table 1 - Ingredients and nutrient compositions of the experimental diets¹ (as-fed basis)

Ingredient	Control	5% SMC	10% SMC	15% SMC
	-----%-----			
Corn	64.25	59.82	55.4	50.82
Soybean meal, 44%	21.5	21.9	22.3	22.8
Spent mushroom compost	0	5.0	10.0	15.0
Wheat bran	5.0	5.0	5.0	5.0
Cane molasses	3.0	3.0	3.0	3.0
Salt	0.3	0.3	0.3	0.3
Dicalcium phosphate	0.8	0.79	0.78	0.77
Limestone, pulverized	1.60	1.62	1.63	1.7
Choline chloride, 50%	0.1	0.1	0.1	0.1
Lysine	0	0.01	0.02	0.03
DL-methionine	0.2	0.21	0.22	0.23
Rice hulls	3.0	2.0	1.0	0
Vitamin premix ²	0.1	0.1	0.1	0.1
Mineral premix ³	0.15	0.15	0.15	0.15
Total	100.0	100.0	100.0	100.0
Calculated values				
Metabolizable energy, kcal/kg	2800	2800	2800	2800
Crude protein, %	15.5	15.5	15.5	15.5
Calcium, %	0.73	0.73	0.73	0.73
Phosphorus, %	0.64	0.64	0.64	0.64

¹ Control: corn-soybean meal; 5%, 10%, and 15% SMC: supplementing 5%, 10%, and 15% of spent mushroom compost (SMC) meal in the basal diet, respectively.

² Provided per kilogram of diet: vitamin A - 10,000 IU; vitamin D3 - 2,000 IU; vitamin E - 20 mg; vitamin K3 - 1.5 mg; vitamin B1 - 1.00 mg; vitamin B2 - 4.8 mg; vitamin B6 - 3.00 mg; vitamin B12 - 16 μg ; folic acid - 0.50 mg; calcium pantothenate - 10.0 mg; niacin - 25 mg; biotin - 2.00 mg.

³ Provided per kilogram of diet: Fe (FeSO_4) - 120 mg; Cu ($\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$) - 22.5 mg; Mn ($\text{MnSO}_4 \cdot \text{H}_2\text{O}$) - 120 mg; Zn (ZnO) - 75.0 mg; I (KI) - 1.275 mg; Co (CoSO_4) - 0.375 mg; Se (Na_2SeO_3) - 0.27 mg.

Results

The proximate composition results showed that SMC has 48.6% dry matter, 16.6% crude protein, and 65.1% crude fiber, respectively. The SMC extracts contained 2.49 ± 0.62 mg gallic acid equivalent/g DW and 1.08 ± 0.15 mg quercetin equivalent/g DW of total phenolics and flavonoid contents.

There was no significant effect between the control and 5% SMC group in body weight ($P < 0.05$) (Table 2). The body weight of the 15% SMC group was significantly lower than that of the control group at the age of 12

Table 2 - Effect of spent mushroom compost on growth performance in grower geese

Item	Treatment ¹				SEM
	Control	5% SMC	10% SMC	15% SMC	
Body weight, kg/bird					
Initial (5 wk)	2.19	2.30	2.19	2.14	0.049
8 wk	4.14ab	4.23a	3.89b	3.89b	0.084
12 wk	4.95a	4.90ab	4.62ab	4.58b	0.085
Feed intake, kg feed/bird					
5-8 wk	7.35	7.21	7.42	7.40	0.140
9-12 wk	6.84	7.01	7.51	7.81	0.322
Feed conversion ratio, kg feed/kg gain					
5-8 wk	3.83	3.82	4.41	4.26	0.219
9-12 wk	8.76	11.5	11.0	14.4	2.220

¹ Results are given as the means of three pens for 10 geese in each pen. Control: corn-soybean meal; 5%, 10%, and 15% SMC: supplementing 5%, 10%, and 15% of spent mushroom compost (SMC) meal in the basal diet, respectively.

SEM - standard error of the mean.

a,b - means without the same letter within the same row under treatment differ significantly ($P < 0.05$).

Table 3 - Effect of spent mushroom compost on serum antioxidant enzyme activities in grower geese at weeks 8 and 12

Item	Treatment ¹				SEM
	Control	5% SMC	10% SMC	15% SMC	
SOD, U/mL					
8 wk	49.4	54.2	54.2	50.2	2.494
12 wk	53.3	51.8	47.5	47.6	4.569
TEAC, U/mL					
8 wk	0.45	0.43	0.45	0.35	0.035
12 wk	0.51	0.49	0.49	0.45	0.021
MDA, U/mL					
8 wk	8.04	7.21	7.35	8.76	1.403
12 wk	8.20a	6.00ab	6.98ab	4.34b	1.004

¹ Results are given as the means of three pens for four geese in each pen. Control: corn-soybean meal; 5%, 10%, and 15% SMC: supplementing 5%, 10%, and 15% of spent mushroom compost (SMC) meal in the basal diet, respectively.

SEM - standard error of the mean.

SOD - superoxide dismutase; TEAC - trolox equivalent antioxidant capacity; MDA - malondialdehyde.

a,b - means within the same rows in the same period but without the same letter are significantly different ($P < 0.05$).

weeks ($P < 0.05$). There were no significant differences among the groups for feed conversion ratio or intake ($P > 0.05$).

There were no significant effects among the groups for serum SOD and TEAC at weeks 8 and 12 ($P > 0.05$). Serum MDA content in the 15% SMC group was lower than that of control group at week 12 ($P < 0.05$; Table 3). There were no differences between control and 5% SMC group for serum MDA at 8 and 12 weeks. The results for the effects of SMC on blood biochemical parameters in grower geese at week 12 reveal no significant effects between the control and SMC groups regarding the blood biochemical parameters in grower geese at week 12 ($P > 0.05$; Table 4).

There were no significant effects between control and SMC groups for carcass variables and meat characteristics of grower geese ($P > 0.05$; Table 5). The color of the meats in the SMC groups had a higher score than that of control group ($P < 0.05$). The flavor and acceptability scores of meats in the 5% SMC group were significantly higher than those of control and 15% SMC groups ($P < 0.05$; Table 6).

Table 4 - Effect of spent mushroom compost on blood biochemical parameters in grower geese at week 12

Item	Treatment ¹				SEM
	Control	5% SMC	10% SMC	15% SMC	
WBC, $10^3/\mu\text{L}$	239	254	219	242	11.59
RBC, $10^6/\mu\text{L}$	1.68	2.01	1.74	1.87	0.117
HB, g/dL	10.8	12.2	10.9	11.4	0.436
HT, %	32.8	36.6	33.0	34.2	1.213
MCV, fl	201	183	195	184	10.73
MCH, pg	66.5	60.8	63.4	61.2	3.418
MCHC, g/dL	33.0	33.3	32.7	33.34	0.314
PLT, $10^3/\mu\text{L}$	25.0	34.8	26.0	26.5	4.374
GLU, mg/dL	142	135	160	144	6.947
BUN, mg/dL	4.00	4.17	4.50	4.33	0.276
UA, mg/dL	2.32	2.05	2.03	1.98	0.265
TP, g/dL	5.83	5.87	6.20	6.92	0.456
ALB, g/dL	2.67	2.68	2.77	2.97	0.162
GLO, g/dL	3.17	3.18	3.43	3.95	0.296
A/G	0.84	0.84	0.81	0.78	0.018
TG, mg/dL	155	139	137	130	15.01
TC, mg/dL	207	215	208	224	13.29
HDL, mg/dL	106	112	106	122	7.624
LDL, mg/dL	72.0	79.0	81.8	81.0	6.746

¹ Results are given as the means of three pens for four geese in each pen. Control: corn-soybean meal; 5%, 10%, and 15% SMC: supplementing 5%, 10%, and 15% of spent mushroom compost (SMC) meal in the basal diet, respectively.

SEM - standard error of the mean.

WBC - white blood cell; RBC - erythrocyte; HB - hemoglobin; HT - hematocrit; MCV - mean corpuscular volume (HT/RBC); MCH - mean corpuscular hemoglobin (HB/RBC); MCHC - mean corpuscular hemoglobin concentration (HB/HT); PLT - platelet; GLU - glucose; BUN - blood urea nitrogen; UA - uric acid; TP - total protein; ALB - albumin; GLO - globulin; A/G - ALB/GLO; TG - triglycerides; TC - total cholesterol; HDL - high-density lipoprotein; LDL - low-density lipoprotein.

Table 5 - Effect of spent mushroom compost on carcass performances and meat characteristic in grower geese at week 12

Item	Treatment ¹				SEM
	Control	5% SMC	10% SMC	15% SMC	
Carcass					
Breast weight, %LW	17.7	17.40	16.8	16.4	0.86
Legs weight, %LW	13.5	13.8	13.4	13.7	0.33
Head and neck weight, %LW	10.4	11.0	10.8	9.96	0.56
Back weight, %LW	14.7	15.9	14.9	16.0	0.59
Paw weight, %LW	3.07	3.26	3.08	3.07	0.18
Abdominal fat pad weight, %LW	2.27	2.21	2.39	2.21	0.28
Liver weight, %LW	1.49	1.59	1.49	1.45	0.10
Gizzard weight, %LW	4.27	3.80	3.93	4.10	0.34
Intestinal weight, %LW	3.57	4.22	4.04	3.94	0.29
Meat characteristic					
Cooking loss, %	32.9	30.5	30.6	29.3	0.881
Shear value, kg/cm ²	7.68	7.13	8.17	8.35	0.793

¹ Results are given as the means of three pens for four geese in each pen. Control: corn-soybean meal; 5%, 10%, and 15% SMC: supplementing 5%, 10%, and 15% of spent mushroom compost (SMC) meal in the basal diet, respectively.

SEM - standard error of the mean.

LW - live weight.

Table 6 - Effect of spent mushroom compost on sensory evaluation in grower geese at week 12

Item	Treatment ¹				SEM
	Control	5% SMC	10% SMC	15% SMC	
Flavor	5.25b	6.94a	5.82ab	5.19b	0.409
Juiciness	6.63a	6.13ab	7.06a	5.25b	0.341
Color	4.38b	6.69a	6.06a	6.44a	0.468
Acceptability	4.81c	7.81a	6.88ab	6.32b	0.502

¹ Results are given as the means of three pens for two geese in each pen. Control: corn-soybean meal; 5%, 10%, and 15% SMC: supplementing 5%, 10%, and 15% of spent mushroom compost (SMC) meal in the basal diet, respectively.

SEM - standard error of the mean.

a,b,c - means without the same letters within the same row under treatment differ significantly (P<0.05).

Discussion

The body weight of the 15% SMC group was significantly lower than that of control group at the age of 5-12 weeks in grower geese, which may suggest that the increased coarse fiber in feed can result in reduced nutrient digestion rate. Foluke et al. (2014) indicated that 75% or 100% SMC replaced with wheat bran in poultry diet led to a significantly lower body weight in starter and finisher phases, as compared with 50% SMC and control groups. The authors suggest that the increased SMC replacement reduced the protein surface digestion rate. Hsu et al. (2000) pointed out that feed with a high content of crude fiber (16%) would reduce the residence time of feed in the stomach and intestines and increase the volume of the stomach and intestines in geese. This may also illustrate why the 15% SMC group had a lower trend

for feed conversion ratio than the 5% SMC group at 9-12 weeks (14.4 vs. 11.5 kg feed/kg gain).

Because free radicals attack the phospholipids of cell membranes, which contain large amounts of unsaturated fatty substances, they may cause lipid peroxidation, be oxidized into lipid peroxide, and finally, be converted to secondary metabolites (such as MDA). The intake of natural phytochemical components (e.g., resveratrol and phenolic components) could effectively reduce the amount of animal serum MDA (Sahin et al., 2010; Lee et al., 2012a). Spent mushroom compost has phytochemical components, such as phenolic compounds and flavonoids, which could significantly protect cells from the damage of free radicals and improve antioxidant capacity (Tuzcu et al., 2008); hence, it could effectively decrease the serum MDA content for the higher SMC-replaced group in this study.

The blood biochemical parameters showed no significant difference when broilers consumed a diet supplemented with 3% or 5% of dietary garlic powder (Choi et al., 2010b; Elagib et al., 2013). Moreover, broilers fed a dietary mixture of medicinal herb extracts (consisting of green tea, garlic, chicory, and cinnamon) showed no influence in the composition of raw thigh meat when compared with control group (Shirzadegan and Falahpour, 2014). Fletcher (2002) showed that the values of shear force increased with age due to an increase in the hardness of connective tissue and in collagen cross-linking, but not in the formula of the feed. This illustrates that there is no significant difference between geese fed SMC and control group in terms of chest carcass performance and meat characteristics.

Kim et al. (2009) pointed out that sensory panelists recorded greater hardness and flavor scores for birds receiving dietary supplementation of 2.0% and 4.0% garlic bulb and husk. Sahoo et al. (2013) used a diet supplemented with 0.05% or 0.1% of an herbal mixture (*Allium sativum*, *Commiphora mukul*, and *Trigonella foenum graecum*) and obtained a lean meat product with increased overall acceptability, as compared with the meat in the control group. In this study, panelists recorded higher flavor levels in meat from geese fed 5% SMC than from the control group. The score for color of meats in the SMC groups was also higher than that of control group. In conclusion, this could suggest better customer acceptability in subsequent sales of products for geese given and 5% SMC in their diet. The 5% corn replaced with 5% SMC meal provides better meat sensory attributes for grower geese; thus, SMC could be utilized in geese diets, and as it is a local resource, it would reduce costs. This strategy can help solve an environmental issue that could result from the accumulation of agricultural wastes.

Conclusions

Spent mushroom compost can be supplemented at 5% in the feed without adversely influencing growth performance or meat characteristics. The level of 5% spent mushroom compost provides better meat sensory attributes for grower geese. Spent mushroom compost can be utilized in geese diets, and as it is a local resource, it may reduce costs.

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