



Feeding Value of Copra Meal in Corn-Animal Protein-Based Diets and Enzyme Supplementation for Egg-Type Birds: Growth Performance Egg, Production and Fatty Acid Profile

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ARTICLE INFO	ABSTRACT
<p><i>Research Article</i></p> <p>Received : 26/09/2020 Accepted : 21/10/2020</p> <p>Keywords: Alternative ingredients Diet composition Enzyme supplementation Egg quality Layer performance</p>	<p>A study investigated the effect of enzyme supplementation of copra meal in corn-animal protein-based diets on pullet growth performance, egg production and fatty acid composition in laying hens. A total of 144, 57 day-old and 20 week-old Shaver Brown pullets were assigned to 8 diets, 2 controls (no copra meal) with and without enzyme and 6 diets containing copra meal at 150, 300 and 450 g/kg with and without enzyme. The experiment was laid as a factorial arrangement (4 copra meal×2 enzyme) in completely randomised design with 3 replicates of 6 birds each per cage for pullet and laying hens. Pullet results showed reduced feed intake on the control diet with enzyme compared to 150, 450 g/kg diets and 300 g/kg diet without enzyme. In the main effects, weight gain was reduced on 300 g/kg copra meal diet. Enzyme supplementation had no effect on growth parameters of pullets. Laying hens results showed significant interaction effects on feed intake and feed conversion ratio but other performance parameters were unaffected by the interaction. Monounsaturated fatty acid of the egg increased on all copra meal diets, saturated fatty acids increased on 450 g/kg copra meal and enzyme supplementation reduced this but interaction had no effect on poly-unsaturated fatty acids. In the main effects, higher egg shape index and deeper yolk colour were observed on 450 g/kg copra meal diet. Inclusion of copra meal at 450 g/kg increased saturated and monounsaturated fatty acid of the egg but copra meal level had no effect on polyunsaturated fatty acid. Enzyme supplementation had no effect on egg parameters reduced saturated fatty acid, increased monounsaturated fatty acid without affecting polyunsaturated fatty acid. In corn-animal protein-based diets, copra meal can be included in pullet and laying hens diets at 450 g/kg without compromising pullet growth performance, egg production and quality.</p>

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Introduction

Traditional feed ingredients for poultry, especially protein sources such as soybeans are not readily available in the South Pacific region. This necessitates research into alternative protein feed ingredients in the region. Copra meal (CM), a by-product of coconut oil extraction, is readily available in the region and has moderate protein content ranging from 150 to 250 g/kg (Devi and Diarra, 2017; Devi et al., 2019). However, high fibre content (420-620 g/kg), mainly in the form of non-starch polysaccharides (NSP) and low amino acid profile (Devi et al., 2019; Sundu et al., 2009; Mael et al., 2019) limit utilisation of CM by poultry. Several dietary manipulations including ingredients selection (Devi and

Diarra, 2017) feed processing methods (Sundu et al., 2009) amino acids and enzyme supplementation (Devi et al., 2019; Sundu et al., 2009; Mael et al., 2019) improved the utilisation CM by poultry (Devi and Diarra, 2017) observed that broiler chickens performed better on CM fed with animal compared to plant protein sources. The residual oil content of CM (35 to 120 g/kg) also makes it a good energy source in poultry diets (Sundu et al., 2009; Devi and Diarra, 2019) but the fat is mainly saturated (Boateng et al., 2016). Although diet composition is known to affect the fatty acid composition of poultry products (Cherian, 2016) the effect of feeding CM on fatty acid composition of poultry egg is limited. This

study investigated the effect of enzyme supplementation of corn-animal protein-based diets containing CM on the performance of laying hens. The following hypotheses were tested:

- i. When fed with animal protein sources, copra meal level will not affect performance of laying hens or fatty acid composition of eggs.
- ii. Supplementation of diets with enzyme will have no beneficial effects.

Materials and Methods

Experimental Site

The investigation consisted of two experiments and was carried out at Ratish Poultry farm, Baulevu road, Nausori, Fiji. The research committee of the University of the South Pacific approved the experimental protocol.

Experimental Protein Sources

The protein sources, fish meal (FM), meat and bone meal (MBM) were purchased locally from Fiji Meat Industries Board (Abattoir) and copra meal (CM) from Fiji Cooperative Dairy Company Limited and analysed for proximate composition, amino acid (AA) and fatty acid profile (Tables 1 and 2). Corn-animal protein-based diets were formulated for each of the experiment.

Experimental Diets

Eight pullet and layer mash diets based on FM and MBM as major protein sources were formulated to contain 15 and 17% CP (crude protein) respectively for the experiment (Table 3 and 4). The diets consisted of 2 controls with no CM with and without enzyme and 6 other diets containing CM at 150, 300 and 450 g/kg with and without enzyme. The enzyme used in this study was Challengzyme 1309A from Beijing Challenge Bio-Technology Company Limited with 8 activities (xylanase 15000 U/g, protease 8000 U/g, pectinase 500 U/g, cellulase 300 U/g, β -glucanase 800 U/g, β -mannanase 100 U/g, α -galactosidase 100 U/g and amylase 500 U/g) (Challengzyme, 2016). Challengzyme was included at 300 g/tonne diet. All diets were fed as mash to the experimental birds.

Experimental Birds and Management

One hundred and sixty day-old Shaver Brown chicks were purchased from Pacific Feeds Limited, Suva, Fiji and warm brooded together on deep litter with wood shavings as litter material for the first 56 days, during which they were fed commercial chick starter feed. On day 57, 144 pullets were weighed and allotted to 24 rearing cages (65.5 cm \times 50 cm \times 35.5 cm) and fed 8 dietary treatments replicated 3 times with 6 birds per replicate in a completely randomized design. The experiment was laid out in a factorial arrangement of 4 levels of CM and 2 levels of enzyme for both pullet and laying hen experiments.

Pullets were vaccinated against Mareks, Infectious Bronchitis and Fowl Cholera at 8, 10 and 12 weeks respectively. On 15th week, a total of 144 point of lay Shaver Brown pullets (140 day-old) were weighed nearest to mean weight (1,206.9 \pm 34.5 g) and transferred to layer cages and allocated to 72 cages with 2 birds per

cage (38.5 cm \times 35 cm \times 40 cm). Feed and water were provided ad-libitum throughout the duration of the experiment to pullets and laying hens for 80 and 105 days respectively. The lighting programme for laying hens was 14 h as per (Shaver management guide, 2016). Stressol was added to drinking water for laying hens to reduce heat stress during the experiment.

Data Collection

Growth Performance

Feed intake (FI) was calculated by difference between the quantity fed and the leftover. Weight gain (WG) was obtained by difference between the initial and final weights. Feed conversion ratio (FCR) was derived as the ratio of feed consumed to weight gained in pullets.

Egg Production and Quality Measurements

Eggs produced were collected and counted per cage and hen-day production (HDP) calculated as:

$$\text{HDP} = (\text{eggs collected}) / (\text{hens present}) \times 100$$

Sample eggs were weighed weekly per cage using a digital scale (Jadever JKH-500 series, Smartfox, Auckland, NZ) sensitive to 0.1 g and egg mass (EM) calculated as:

$$\text{EM} = \text{eggs collected} \times \text{mean egg weight (g)}$$

Feed conversion ratio (FCR) was calculated as the ratio of unit feed consumed to unit egg produced as:

$$\text{FCR} = \text{feed consumed} / \text{eggs mass}$$

Sample eggs from each treatment were used for egg quality measurements on a weekly basis. Egg weight was taken using a digital scale sensitive to 0.1 g. Egg length and width were measured using a digital Vernier caliper and shape index (SI) calculated as:

$$\text{SI} = \text{Egg width} / \text{egg length} \times 100$$

Eggs were then broken on a glass surface to separate the yolk carefully from the albumin. Yolk colour was determined using a Roche yolk colour fan. Albumin height was measured using a tripod spherometer. Shell thickness was taken as the mean of 3 measurements (broad end, middle and narrow end) using a digital Vernier caliper. Assessment of egg freshness was based on the Haugh unit (HU). Haugh unit (HU) was calculated according to Raymond Haugh (1937) cited in Eisen et al., 1962) as:

$$\text{HU} = 100 \times \log (h - 1.7w^{0.37} + 7.6)$$

Where h=height of the albumen and w=weight of the egg. At the end of the experiment, a total of 32 eggs (4 eggs per treatment) were randomly selected for fatty acid composition at the Institute of Applied Sciences laboratory, University of the South Pacific, Laucala Campus, Suva.

Data Analysis

Proximate analysis (dry matter, crude protein, crude fibre and fat) was done according to (AOAC, 1990) procedures. Amino acid profile was determined using the Performic acid oxidation method (AOAC, 1997). Total fat was determined according to AOAC (2012) (ID 922.06) modified method and the fatty acid profiling done using

Hewlett Packard 6890® gas chromatograph (Sukhija and Palmquist, 1988). Total (TDF), soluble (SDF) and insoluble detergent fibre (IDF) was analysed using Megazyme as per AOAC 991.45. Neutral detergent fibre (NDF) and acid detergent fibre (ADF) were analysed using Tecator Fibertec according to AOAC 2002.04 and AOAC 973.18, respectively.

Table 1. Composition of the experimental protein sources in selected constituents

Constituents (g/kg)	Protein sources		
	FM	MBM	CM
ME MJ/kg	11.31	13.44	10.88
Dry matter	725	879	887
Crude protein	531	481	184
Ether extract	106	253	120
Ash	117	144	49
IDF	-	-	402
SDF	-	-	17
TDF	-	-	419
Crude fibre	5	21	189
NDF	222	250	441
ADF	26	82	271

ME: metabolisable energy; FM: fish meal; MBM: meat and bone meal; CM: copra meal; IDF: insoluble dietary fibre; SDF: soluble dietary fibre TDF: total dietary fibre; NDF: neutral detergent fibre; ADF: acid detergent fibre

Table 2. Fatty acid composition and amino acid profile of the experimental protein sources

Fatty acids (g/100g DM)	Systematic name	Formulae	FM	MBM	CM
C _{6:0} Caproic	Hexanoic acid	C ₆ H ₁₂ O ₂	ND	0.01	0.12
C _{8:0} Caprylic	Octanoic acid	C ₈ H ₁₆ O ₂	ND	0.01	1.26
C _{10:0} Capric	Decanoic acid	C ₁₀ H ₂₀ O ₂	<0.01	0.02	0.89
C _{11:0} Undecylic	Undecanoic acid	C ₁₁ H ₂₂ O ₂	<0.01	<0.01	<0.01
C _{12:0} Lauric	Dodecanoic acid	C ₁₂ H ₂₄ O ₂	<0.01	0.02	6.41
C _{13:0} Tridecylic	Tridecanoic acid	C ₁₃ H ₂₆ O ₂	<0.01	0.01	0.01
C _{14:0} Myristic	Tetradecanoic and	C ₁₄ H ₂₈ O ₂	0.21	0.69	2.43
C _{16:0} Palmitic	Hexadecanoic acid	C ₁₆ H ₃₂ O ₂	1.93	5.99	1.21
C _{17:0} Margaric	Heptadecanoic acid	C ₁₇ H ₃₄ O ₂	0.13	0.43	0.01
C _{18:0-3} Stearic	Octadecanoic acid	C ₁₈ H ₃₆ O ₂	2.07	12.66	1.55
C _{20:1-5} Arachidic	Eicosanoic acid	C ₂₀ H ₄₀ O ₂	0.77	0.52	0.01
C _{21:0} Heneicosylic	Heneicosanoic acid	C ₂₁ H ₄₂ O ₂	0.02	0.04	0.02
C _{22:0-6} Behenic	Docosanoic acid	C ₂₂ H ₄₄ O ₂	2.3	1.19	0.01
C _{23:0} Tricosylic	Tricosanoic acid	C ₂₃ H ₄₆ O ₂	0.01	0.01	ND
C _{24:0-1} Lignoceric	Tetracosanoic acid	C ₂₄ H ₄₈ O ₂	0.1	0.07	0.02
Amino acids (mg/100mg DM)					
Aspartic acid	L-Aspartic acid	C ₄ H ₇ NO ₄	4.58	3.61	1.48
Threonine	L-Threonine	C ₄ H ₉ NO ₃	2.16	1.67	0.56
Serine	L-Serine	C ₃ H ₇ NO ₃	2.13	1.90	0.79
Glutamic acid	L-glutamic acid	C ₅ H ₉ NO ₄	6.68	5.63	3.12
Proline	L-Proline	C ₅ H ₉ NO ₂	2.84	3.20	0.62
Glycine	2-Aminoethanoic acid	C ₂ H ₅ NO ₂	4.61	5.21	0.81
Alanine	L-Alanine	C ₃ H ₇ NO ₂	3.44	3.16	0.80
Valine	L-Valine	C ₅ H ₁₁ NO ₂	2.27	1.95	0.89
Isoleucine	L-Isoleucine	C ₆ H ₁₃ NO ₂	1.85	1.45	0.56
Leucine	L-leucine	C ₆ H ₁₃ NO ₂	3.46	2.87	1.10
Tyrosine	L-tyrosine	C ₉ H ₁₁ NO ₃	1.50	1.18	0.47
Phenylalanine	L-Phenylalanine	C ₉ H ₁₁ NO ₂	1.87	1.56	0.74
Histidine	L-Histidine	C ₆ H ₉ N ₃ O ₂	1.50	1.26	0.39
Lysine	L-Lysine	C ₆ H ₁₄ N ₂ O ₂	3.84	2.75	0.64
Arginine	L-arginine	C ₆ H ₁₄ N ₄ O ₂	3.29	3.12	2.35
Cysteine	L-Cysteine	C ₃ H ₇ NO ₂ S	0.36	0.25	0.28
Methionine	DL-methionine	C ₅ H ₁₁ NO ₂ S	1.41	0.90	0.31
Tryptophan	L-Tryptophan	C ₁₁ H ₁₂ N ₂ O ₂	0.46	0.30	0.15

FM: fish meal; MBM: meat and bone meal; CM: copra meal; DM: dry matter; ND: not detected

Table 3. Ingredient composition and calculated analysis of pullet grower diets

Ingredients(g/kg)	CM (g/kg) with no enzyme				CM (g/kg) with enzyme			
	0	150	300	450	0	150	300	450
Corn	543.9	463.5	382.9	302.3	543.7	463.3	382.7	302.1
Wheat middling	272	231.7	191.5	151.2	271.9	231.6	191.3	151
Tuna fish meal	42.4	32.5	22.6	12.7	42.4	32.5	22.6	12.7
Meat & bone meal	84.9	65.1	45.2	25.4	84.9	65.1	45.3	25.5
Copra meal	0	150	300	450	0	150	300	450
Sand	30	30	30	30	30	30	30	30
Limestone	20	20	20	20	20	20	20	20
*Premix	2.5	2.5	2.5	2.5	2.5	2.5	2.5	2.5
Lysine	0.8	1	1.5	2	0.8	1	1.5	2
Methionine	0.5	0.7	0.8	0.9	0.5	0.7	0.8	0.9
Enzyme	0	0	0	0	0.3	0.3	0.3	0.3
Salt	3	3	3	3	3	3	3	3
Calculated analysis								
Crude protein (%)	15	15	15	15	15	15	15	15
ME (MJ/kg)	11.9	11.8	11.7	11.6	11.9	11.8	11.7	11.6
Lysine (g/kg)	7.9	7.7	7.6	7.6	7.9	7.7	7.6	7.6
Methionine (g/kg)	3.5	3.6	3.7	3.8	3.5	3.6	3.7	3.8

*Premix (Vitamin and mineral) Bio-mix supplied/kg diet, vitamin A: 10 000 IU, vitamin D3: 2000 IU, vitamin E: 23 mg, niacin: 27.5mg, vitamin B1: 1.8 mg, B2: 5mg, B6: 3mg, B12: 0.015mg, vitamin K: 3.2mg, pantothenic acid:7.7mg, biotin:0.06mg, folic acid: 0.75mg, choline chloride: 300mg, cobalt: 0.2mg, copper: 3mg, iodine: 1mg, iron: 20mg, manganese: 40mg, selenium: 0.2mg, zinc: 30mg, anti-oxidant: 1.25mg; CM: Copra meal; ME: Metabolisable Energy

Table 4. Ingredient composition and calculated analysis of the layer diets

Ingredients (g/kg)	CM (g/kg) with no enzyme				CM (g/kg) with enzyme			
	0	150	300	450	0	150	300	450
Corn	499.2	421	342.1	263.4	499.1	420.5	341.9	263.2
Wheat middling	249.6	210.1	171	131.7	249.4	210.3	170.9	131.6
Tuna fish meal	51.5	40.6	29.7	18.8	51.5	40.6	29.7	18.8
Meat & bone meal	102.9	81.1	59.4	37.7	102.9	81.1	59.4	37.7
Copra meal	0	150	300	450	0	150	300	450
Sand	40	40	40	40	40	40	40	40
Limestone	50	50	50	50	50	50	50	50
*Premix	2.5	2.5	2.5	2.5	2.5	2.5	2.5	2.5
Lysine	0.8	1	1.5	2	0.8	1	1.5	2
Methionine	0.5	0.7	0.8	0.9	0.5	0.7	0.8	0.9
Enzyme	0	0	0	0	0.3	0.3	0.3	0.3
Salt	3	3	3	3	3	3	3	3
Calculated analysis								
Crude protein (%)	17	17	17	17	17	17	17	17
ME (MJ/kg)	12.2	12	11.8	11.5	12.2	12	11.7	11.5
Lysine (g/kg)	8.7	8.2	8.1	8.0	8.7	8.2	8.1	8.0
Methionine (g/kg)	3.6	3.8	3.8	3.9	3.6	3.8	3.8	3.9
Calcium (g/kg)	32.8	30.1	27.4	24.7	32.8	30.1	27.4	24.7
Phosphorus (g/kg)	10.7	9.5	8.3	7.1	10.7	9.5	8.3	7.1
Ca:P	3.1:1	3.2:1	3.3:1	3.5:1	3.1:1	3.2:1	3.3:1	3.5:1

*Premix (Vitamin and mineral) Bio-mix supplied/kg diet, vitamin A: 10 000 IU, vitamin D3: 2000 IU, vitamin E: 23mg, niacin: 27.5mg, vitamin B1: 1.8 mg, B2: 5mg, B6: 3mg, B12: 0.015mg, vitamin K: 3.2mg, pantothenic acid:7.7mg, biotin:0.06mg, folic acid: 0.75mg, choline chloride: 300mg, cobalt: 0.2mg, copper: 3mg, iodine: 1mg, iron: 20mg, manganese: 40mg, selenium: 0.2mg, zinc: 30mg, anti-oxidant: 1.25mg; CM: Copra meal; ME: Metabolisable energy.

Gross energy (solid) was determined in the Bomb calorimeter and Metabolisable energy (ME) calculated according to Fisher and Boorman (1986) as:

$$ME (Kcal/Kg) = 37 \times CP + 81 \times EE + 35.5 \times NFE.$$

Where CP=crude protein; NFE=nitrogen-free extract and EE=ether extract.

Statistical Analysis

Data collected were subjected to ANOVA (Steel and Torrie, 1980) of a factorial arrangement using the GLM of SPSS (Windows, version 22.0; IBM Corp, Armonk, NY, USA) (SPSS, 2013). Individual were the experimental units for weight change whereas cages were the experimental units for feed intake.

Treatment means were compared using the Least Significant Difference (LSD) and differences considered significant when ($P < 0.05$).

Results

Pullet Growth Performances

From the growth performance results of pullets (Table 5) WG and FCR were not affected by the interaction of CM and enzyme but there was a significant interaction effect on FI ($P<0.05$). Enzyme supplementation reduced the intake of the control diet compared to 150 and 450 g/kg and 300 g/kg CM without enzyme ($P<0.05$).

There was no significant interaction effect on FI between the negative control and CM based diets ($P>0.05$). In the main effect, weight gain was depressed on 300 g/kg CM diet ($P<0.05$). A better FCR was observed on control compared to the CM diets ($P<0.05$). Enzyme addition did not affect any of the performance parameters studied ($P>0.05$). No mortality was recorded throughout the pullet experiment.

Laying Performance

The egg performance results are presented in Table 6. No mortality was recorded throughout layer experiment. Feed intake and FCR showed significant interaction effects ($P<0.05$) but egg mass remained unaffected ($P>0.05$). Feed intake significantly reduced on the 300 g CM/kg with enzyme compared to the other CM-based diets ($P<0.05$). Feed conversion ratio was poorer on 300 g CM without enzyme and 450g CM with enzyme ($P<0.05$) compared to control diets and 150g CM without enzyme.

In the main effects, CM level had no effects on FI, HDP, MEW, EM, HU and shell thickness ($P>0.05$). Feed conversion ratio increased above 150 g CM/kg ($P<0.05$). Lower shape index was observed on 300 g CM/kg compared to 450 g CM/kg ($P<0.05$). A deeper yolk colour was recorded on the 450 g CM/kg ($P<0.05$) but yolk colour did not differ among the control, 150 and 300 g CM/kg ($P>0.05$). Enzyme supplementation had no effects on any of the performance traits evaluated ($P>0.05$).

Fatty Acid Composition of Eggs

The fatty acids SFA (saturated), MUFA (monounsaturated) and PUFA (polyunsaturated) compositions (g/100g) of the eggs are presented in Table 7. Saturated fatty acid and MUFA composition was significantly affected by the interaction ($P<0.05$) but PUFA was unaffected ($P>0.05$). Lower values of SFA were recorded on the control and the 300 g CM with and without enzyme and higher MUFA on the 450g CM with enzyme ($P<0.05$). In the main effects, higher values of SFA and MUFA were observed on 450 g/kg CM ($P<0.05$) but PUFA was not affected CM level ($P>0.05$). Enzyme supplementation reduced SFA and increased MUFA ($P<0.05$) but had no effect on PUFA composition ($P>0.05$).

Table 5. Growth performances of pullets fed increasing CM levels in corn-animal protein-based diets and Challengezyme supplementation

Treatment		FI (kg/cage)	WG (kg/cage)	FCR
Copra	Enzyme			
0	No	24.44 ^{ab}	3.84	6.45
	Yes	23.16 ^b	4.44	5.22
150	No	25.42 ^a	4.21	6.07
	Yes	26.16 ^a	3.77	6.95
300	No	26.45 ^a	3.45	7.68
	Yes	24.23 ^{ab}	3.71	6.68
450	No	25.28 ^a	4.05	6.29
	Yes	27.11 ^a	4.29	6.51
SEM		0.691	0.269	0.534
Main effects				
Copra				
0		23.80	4.14 ^a	5.84 ^b
150		25.79	3.99 ^a	6.51 ^a
300		25.34	3.58 ^b	7.18 ^a
450		26.20	4.17 ^a	6.40 ^a
Enzyme				
No		25.40	3.89	6.62
Yes		25.17	4.05	6.34
Probabilities				
Enzyme		0.057	0.048	0.035
Copra		0.644	0.401	0.466
Copra*Enzyme		0.038	0.306	0.195

FCR: feed conversion ratio; FI: feed intake; WG: weight gain; SEM: standard error of mean; a, b, c: values within the column with different superscripts differ significantly ($P<0.05$).

Table 6. Performance of laying hens fed varying CM levels in corn-animal protein diets and Challengzyme supplementation

Treatment		FI	HDP	MEW	EM	FCR	HU	SI	YC	ST
Copra	Enzyme	(kg)	(%)	(g)	(kg)	(FI:EM)				(mm)
0	No	35.07 ^{ab}	73.9	48.3	12.97	2.73 ^c	80.6	77.4	5.8	43
	Yes	36.26 ^a	75.2	46.5	13.10	2.78 ^c	75.8	78.0	5.8	46
150	No	36.19 ^a	81.9	47.2	13.61	2.66 ^c	79.9	78.3	5.0	41
	Yes	38.43 ^a	72.6	47.7	12.00	3.24 ^b	81.0	77.2	5.0	41
300	No	38.68 ^a	68.2	47.1	11.37	3.42 ^a	83.4	75.9	5.8	44
	Yes	34.92 ^b	72.7	48.5	12.17	2.88 ^{bc}	82.1	76.5	5.0	41
450	No	37.26 ^a	74.7	50.2	13.44	2.80 ^{bc}	80.2	78.9	7.5	44
	Yes	37.49 ^a	65.6	47.4	11.07	3.44 ^a	86.5	78.8	5.8	42
SEM		0.923	3.721	1.347	0.828	0.169	2.687	0.702	0.589	0.018
Main effects										
Copra										
0		35.66	74.5	47.4	13.04	2.75 ^b	78.2	77.7 ^{ab}	5.8 ^b	44
150		37.31	77.3	47.5	12.81	2.95 ^{ab}	80.5	77.8 ^{ab}	5.0 ^b	41
300		36.80	70.5	47.8	11.77	3.15 ^a	82.8	76.2 ^b	5.4 ^b	43
450		37.38	70.2	48.8	12.26	3.12 ^a	83.4	78.8 ^a	6.7 ^a	43
Enzyme										
No		36.80	74.7	48.2	12.85	2.90	81.0	77.6	6.0	43
Yes		36.77	71.6	47.5	12.09	3.08	81.4	77.6	5.4	43
Probabilities										
Copra		0.258	0.209	0.706	0.441	0.013	0.242	0.015	0.046	0.407
Enzyme		0.967	0.250	0.480	0.210	0.149	0.864	0.979	0.153	0.797
Copra*Enzyme		0.026	0.182	0.402	0.231	0.010	0.254	0.578	0.455	0.395

FI: feed intake; EM: egg mass; HDP: hen day production; MEW: mean egg weight; FCR: feed conversion ratio; HU: Haugh Unit; SI: Shape index; YC: Yolk colour; ST: Shell Thickness SEM: standard error of mean; a, b, c: values within the column with different superscripts differ significantly (P<0.05)

Discussion

Chemical Analysis

The CP content of the experimental CM (184 g/kg) is comparable to 180-210 g/kg reported by (Sundu et al., 2014; Diarra et al., 2018). Other authors (Devi and Diarra, 2017; Sundu et al., 2009; Sundu et al., 2006; Diarra et al., 2014) reported up to 250 g/kg CP in CM. The AA composition showed slightly higher profiles of lysine (6.4 g/kg) and methionine (3.1 g/kg) than the values (4.7 and 2.9 g/kg, respectively) reported by (Devi and Diarra, 2017). The experimental CM contained about 420 g/kg total NSP. (Knudsen, 1997) also reported same value (420 g/kg) of NSP in CM. However, other authors (Devi and Diarra, 2017; Devi et al., 2019) reported higher NSP (520-610 g/kg) in CM. Several factors including coconut variety, harvest age, drying process, oil extraction method and storage conditions affect the composition of CM (Sundu et al., 2006; Diarra et al., 2014; Panigrahi, 1992). The fatty acid composition of the experimental CM characterised by higher SFA and lower unsaturated fatty acids is in agreement with earlier reports (NRC, 1994; Boateng et al., 2016).

Pullet Growth Performance

Feed intake was reduced on the control diet with enzyme compared to 150 and 450 g/kg diets and 300 g/kg CM without enzyme. This may be attributed to increased nutrient availability resulting from enhanced hydrolysis of the low fibre control diet suggesting that the enzyme concentration was not sufficient enough to hydrolyse CM at these inclusion levels. Diarra et al. (2018) also reported higher FI in 56-132 days' old growing pullets fed 200 g CM /kg with enzyme compared to control and attributed

this to faster digesta transit in the gastro intestinal tract (GIT). The higher intake of the CM diets may be attributed to several factors including i) an attempt by the birds to meet their nutrient need, and ii) the beneficial effects of fibre on GIT development (Mateos et al., 2012) suggested that before the onset of egg production, dietary fibre improves GIT development in birds. Bouvarel and Nys (2013) and Pottguetter (2015) also confirmed that dietary CF is needed for stimulation of GIT development and FI at starter phase. In the main effect, the reduced WG on 300 g CM /kg diet compared to other CM diets despite similar FI was not understood. The results of this study are in agreement with those of Diarra et al. (2018) who found reduced WG in pullets fed 200 g/kg CM. Moorthy and Viswanathan; 2006; Moorthy and Viswanathan; 2010) also reported reduced WG and egg production of laying hens above 150-200 g/kg CM in the diet. Contrary to the findings of this study (Sundu et al., 2005) found improved WG in 6 weeks old Ross male broilers fed 300 g/kg CM with enzyme (Hemicel, mannanase®, β -mannanase and Allzyme SSF®) compared to diets without enzyme. The class of poultry, diet composition, enzyme source and concentration are all possible reasons for the differences in the utilisation of CM by poultry.

The improved WG on 450 g/kg compared to 300 g/kg CM diet in this study was not clear but possible enhanced gut health on this fibrous diet may be speculated. The poor FCR on CM diets compared to control diets was mainly attributed to the linear increase in FI on CM based diets compared to the control (Diarra et al., 2014; Diarra et al. (2015) also reported poor FCR in pullets fed CM diet at 200 g/kg without enzyme.

Table 7. Fatty acid compositions SFA, MUFA, PUFA (g/100g) of eggs of laying hens fed increasing CM levels in corn-animal protein-based diets with and without Challengzyme

Treatment		SFA	MUFA	PUFA
Copra	Enzyme			
0	No	2.77 ^c	1.07 ^b	0.095
	Yes	3.60 ^b	0.80 ^b	0.096
150	No	3.70 ^b	0.87 ^b	0.091
	Yes	3.63 ^b	0.83 ^b	0.093
300	No	2.57 ^c	0.80 ^b	0.092
	Yes	2.67 ^c	0.73 ^b	0.091
450	No	4.80 ^a	0.97 ^b	0.095
	Yes	3.87 ^b	2.17 ^a	0.097
SEM		0.152	0.117	0.009
Main effects				
Copra				
0		3.18 ^c	0.93 ^b	0.094
150		3.67 ^b	0.85 ^b	0.092
300		2.62 ^d	0.77 ^b	0.091
450		4.33 ^a	1.57 ^a	0.096
Enzyme				
No		3.46 ^a	0.93 ^b	0.095
Yes		3.44 ^b	1.13 ^a	0.092
Probabilities				
Copra		0.001	0.001	0.560
Enzyme		0.030	0.022	0.560
Copra*Enzyme		0.001	0.001	0.560

SFA: saturated fatty acids; MUFA: monounsaturated fatty acids; PUFA: polyunsaturated fatty acids; SEM: standard error of mean; a,b,c: values within the column with different superscripts differ significantly (P<0.05).

Laying Performance

Egg performance parameters were not affected by the interaction of CM and enzyme probably due to the similarity in nutrient content of the experimental diets and the lower requirements of egg-type birds. These findings are consistent with those reported by earlier researchers (Panigrahi, 1989) reported no adverse effect of feeding 100 and 200 g/kg CM in Star Cross 288 laying hens. Diarra et al. (2014) reported similar HDP on diets containing 200 g CM /kg with or without enzyme and control commercial diet in Shaver Brown laying hens. Abeysekara and Atapattu, (2016) also observed no effect of 100 g CM /kg with enzyme in corn-soybean meal based diets on egg production in 13 week-old Japanese quails. Contrary to our results however Moorthy and Viswanathan, (2010) reported lower HDP on 200 g CM /kg in SBM compared to FM-based diets in 21 to 52 week-old single comb white leghorn (SCWL) layers. Dairo and Fasuyi (2008) found that fermented CM diets at 86.5 and 173 g/kg produced higher HDP compared to control diet in 37 week-old Black Australorp laying hens. The trend of laying performance in the present study could probably be attributed to possible longer digesta retention of CM-based diets resulting in increased nutrient absorption in the GIT. Diarra et al. (2014) attributed improved laying performance and heavier eggs of hens to longer digesta retention and increased absorption of nutrients. In contrast, Diarra et al. (2018) found no improvement in egg performance of Shaver Brown pullets fed 200 g/kg CM in corn-FM-SBM based diets supplemented with Allzyme SSF (0.3 g/kg) compared to CM diets without enzyme Abeysekara and Atapattu (2016) also found no effect of 0.1 g/kg Bio Grain CG enzyme in 200 g/kg CM on egg parameters of Japanese quails. The

performance differences among studies could be attributed to several factors including the species, breed, age of birds, CM processing, composition of basal diet, enzyme source and concentration. The higher value of egg shape index on 450 g/kg CM is in agreement with earlier reports. Dairo and Fasuyi (2008) reported higher shape index on 500 g/kg fermented CM diet in corn-SBM based diets in 37 week-old Black Australorp laying hens. Krawczyk et al. (2013) found significant effect of feeding diets based on oil seed meals and distillers spent grain on egg shape index in 18 week-old White Leghorn and Rhode Island Red laying hens. The authors also observed lighter yolk colour of eggs on wheat-corn-SBM based diet compared to pigment supplemented control and attributed this to lower xanthophyll and carotenoid content of wheat. Panaitea et al. (2019) observed darker yolk colour score on 75 g/kg dried tomato waste in corn-SBM diets compared to control in 96 Tetra SL laying hens and attributed this to the higher carotenoid content of dried tomato waste. The improvement in yolk colour on 450 g CM/kg in this study was not clear but possibly, due to enhancement in carotenoid absorption by the residual fat in CM. Gül et al. (2012) found darker yolk colour in Hisex Brown laying hens fed corn-SBM based diets supplemented with canola oil (20, 40 and 60 g/kg) compared to control.

Fatty Acid Composition of Eggs

Egg yolks of the groups fed 450 g/kg CM contained higher values of SFA and MUFA. The higher inclusion of CM and high intake of this diet may explain the pattern of fatty acid in egg yolks because poultry have limited ability to transform dietary fat. Gül et al. (2012) observed higher egg yolk MUFA in Hisex Brown laying hens fed corn-

SBM based diets with canola oil (20, 40 and 60 g/kg) compared to the control without oil. Recently, Panaitea et al. (2019) found higher egg yolk omega 3 (PUFA) in 96 Tetra SL laying hens fed corn-SBM diets containing 50 g/kg dried tomato waste with flaxseed. The authors attributed this to the fatty acid composition of flaxseed and dried tomato waste in the diet and the anti-oxidant property of lycopene content of tomato waste. In an earlier report Cherian and Sim, (1991) also found increased omega 3 fatty acid of flax and canola seeds in eggs compared to control. Panaitea et al. (2019) observed significant effect of diet on fatty acid profile of egg yolk of White Leghorn and Rhode Island Red laying hens fed oil seed meals and distillers spent grain in wheat-corn-SBM based diets. Contrary to these findings Farias et al. (2019) found no effect of 250 g/kg stored CM corn-SBM diets on egg fatty acid composition of Japanese quails. This suggests that CM quality is an important factor affecting its fatty acid composition. The lower SFA and MUFA despite high FI on 300 g/kg CM diet with enzyme was not be clear and needs further investigation.

Conclusions

Based on the results, it is concluded that CM can be included up to 450 g/kg in corn-animal protein based diets for pullets and laying hens without compromising pullet growth performance, egg production and quality. Inclusion at 300 and 450 g/kg CM increases FCR in laying hens. Enzyme supplementation reduced egg SFA and increased MUFA composition of eggs. Saturated fatty acids of eggs can be altered at higher CM inclusion with enzyme. Copra meal inclusion will have cost advantage and add value to CM, where it is readily available. More research into the source of CM, composition of basal diet, enzyme source and concentration above 30% CM is recommended.

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Conflict of Interest

The authors declare no conflict of interest

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