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Effects of graded levels of rumen digesta based diets with or without enzyme supplementation on the blood chemistry of weaner rabbits

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Abstract

Eight weeks feeding experiment was conducted to investigate the effect of replacing maize with Dried Rumen Digesta (DRD) with or without enzyme supplementation on the haematology and serum chemistry of weaner rabbits. Thirty six weaner rabbits between 4 and 5 weeks old were randomly allocated to six dietary treatments of six rabbits per treatment in a Completely Randomised Design (CRD). There were three replicates of two rabbits each. DRD replaced maize at 0%, 12.5%, 25% levels. There was no significant difference ($P>0.05$) among the WBC values of the experimental animals and the values were within the normal range for Rabbits. There were significant differences ($P<0.05$) in the RBC and the PVC among the treatments. The Mean Corpuscular Haemoglobin Concentration (MCHC) values of T3 and T5 showed no significant differences ($P>0.05$). The result also showed significant differences ($P<0.05$) between T2 and the rest diets. The protein concentration, Cholesterol, ALT, AST and potassium levels showed no significant differences ($P<0.05$) among the treatments. In conclusion, the study suggested up to 25% DRD replacement.

Keywords: Haematology, serology, rabbit, rumen digesta, enzyme

Introduction

Rabbit is a herbivorous, monogastric and pseudo-ruminant animal that can effectively and efficiently convert fodder to meat (Leba, 1980) ^[1]. Rabbit is blessed with unique digestive tract that can convert fibrous materials to animal protein. One of the challenges of animal researchers and nutritionists, is to provide feeding strategies for monogastric animals especially poultry pigs and rabbits. The strategies should be able to minimize the cost of production and reduce competition between animal and human for the feed ingredients (Biobaku, 2002) ^[5].

The nutrition of the rabbits is one of the most important aspects of its production. The non-availability and/or insufficient supply of rabbit pellets which consist mainly of cereal grains hampered their production (Aduku and Olukosi, 1990) ^[2]. The search for alternative sources of energy in rabbit remained a topical issue among animal nutritionists (Igwebuikwe, 2001) ^[9]. The need to improve rabbit production in Nigeria to increase supplies of animal protein is clear, due to the high cost of beef and chicken, also the animal protein shortage facing Nigeria cannot be solved by large animals with their slow production cycle. Animal like rabbit, which are prolific and has short gestation period, can help this protein shortage problem. Rabbit can be produced on forages alone although production can improve by adding other feed supplements.

Rabbit possesses various attributes that are advantageous in comparison to other livestock. Rabbit meat is of excellent protein quality, low in total as well as saturated fat, cholesterol and sodium, therefore, rabbit production is considered as a good source of meat in tropical developing countries where there is abundance of by-products which can be used as feedstuff. Currently commercial rabbits are reared mostly on expensive poultry feeds, as commercial feed are available for rabbit but not widely used because it is expensive. This is one of the important factors limiting the expansion of rabbit farming in Nigeria and other developing nations.

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The rumen content is obtained from the abattoir and cannot be consumed by man. Rumen content is made up of undigested feed eaten by the ruminants animal with lots of micro-organism that aid in feed degradation, the feed and synthesized protein, fatty acids and vitamins produced by the microbes. Rumen content is highly fibrous, hence supplements that can aid digestion of fiber such as enzyme may be needed when using rumen content in feed formulation especially for monogastric animals. When rumen content (R C) is processed to feedstuff, it has repulsive and inherent colour which affects the acceptability when fed to livestock (Balogun *et al.*, 2001) [4].

Enzymes are one of the many types of protein in biological systems. Their essential characteristic is to catalyze the rate of reaction but without being altered by it. Several studies carried out on poultry and pigs have shown that the poor quality feed can be enhanced by supplementation or supplemental enzymes. There is increase in the demand for cereal gains for human consumption, industrial use and as a major ingredient in livestock feed, thus making it unaffordable for small scale farmers. Hence the search for alternative energy sources that could be of value and cheaper.

Materials and methods

Thirty six (36) mix-bred, 4-5 weeks weaner rabbits with a balanced initial determined weight were used. The animals were randomly allocated to six dietary treatment groups in a Completely Randomized Design (CRD). The experiment was replicated thrice with two rabbits per replicate thereby giving a total of six rabbits per treatment. Rumen content was collected fresh from the abattoir immediately the visceral of the cattle was opened and boiled for about two hours with constant stirring to prevent burning. After boiling, it was sun-dried to reduce the moisture content to about 12%. The sun-dried material was subjected to particle size reduction followed by proximate analysis.

Six diets were compounded for the experiment. The ingredients used for formulating the diet include G.N.C, maize, wheat offal, bone meal common salt, vitamin premix, lysine methionine and rumen digesta. The experimental ingredients was included at 0% DRD without enzyme (T1), 0% DRD with enzyme (T2), 12.5% DRD without enzyme (T3), 12.5% DRD with enzyme (T4), 25% DRD without enzyme (T5) and 25% DRD with enzyme (T6) respectively. The (DRD) replaced maize weigh for weight each diet was formulated to meet the minimum protein requirement (16 – 18%CP) of weaner rabbits.

Within the last week of the feeding trial, one rabbit from each replicate was selected at random. The bleeding was done in the morning before feeding. 10ml of blood was obtained from the jugular vein using a sterilized needle and syringe into a sample bottle. EDTA bottle containing anticoagulant and another set of plain bottles without anticoagulants were used for this exercise. About 3ml of the blood sample was put into the EDTA sample bottle while the remaining was put into the plain sample bottle. The samples in the plain bottles were allowed to clot so as to obtain the serum that was used in the determination of some serum metabolites as described by Toro and Ackermann (1975) [17] and Kaneko (1989) [10].

Parameters evaluated include: Red Blood Cell counts (RBC), Parked Cell Volume (PCV), Haemoglobin (HB), Mean Corpuscular Volume (MCV), Mean Corpuscular Haemoglobin (MCH), Mean Corpuscular Haemoglobin Concentration (MCHC) and White Blood Cell counts (WBC). Serological parameters evaluated are total protein, glucose, urea and cholesterol.

Analytical techniques

Packed cell volume was determined by microhematocrit method (Igene and Iboh, 2004) [7]. Haemoglobin content was determined as described by Jain (1993) [8] and Red blood cell was measured with the aid of Neubaur counter (haemocytometer) as reported by Oni *et al.* (2010) [13]. Total protein determination was carried out by the Biuret method of Savory and Sunderman (1968) [16], serum electrolytes was determined by atomic absorption spectroscopy as reported by Adejumo and Onifade (2005) [11].

Data Analysis

The data generated were subjected to analysis of variance (ANOVA) using the SAS package (1999-2000). Fisher LSD was used to separate means ($P < 0.05$) where differences manifest.

Table 1: Percentage composition of experimental diets Brewers Dried Grain Inclusion Levels

Ingredient	0%		12.5%		25 %	
Maize	48	48	42	42	36	36
Wheat offal	25	25	25	25	25	25
Rumen digesta	0	0	12.5	12.5	25	25
Groundnut cake	21	21	21	21	21	21
Bone meal	2.8	2.8	2.8	2.8	2.8	2.8
Salt	0.35	0.35	0.35	0.35	0.35	0.35
Vitamin premix	0.25	0.25	0.25	0.25	0.25	0.25
Lysine	0.3	0.3	0.3	0.3	0.3	0.3
Methionine	0.3	0.3	0.3	0.3	0.3	0.3
Limestone	2	2	2	2	2	2
Total	100	100	100	100	100	100

Calculated

CP (%)	18.1	18.1	18.7	18.7	19.3	19.3
ME (K Cal/kg)	2676	2676	2614	2614	2551	2551
EE (%)	3.76	3.76	4.00	4.00	4.24	4.24
CF%	5.07	5.07	5.59	5.59	6.12	6.12
Ca%	1.53	1.53	1.55	1.55	1.56	1.56
P%	0.54	0.54	0.55	0.55	0.55	0.55

Results and discussion

Results of the effect of the dietary treatments on the hematological parameters and serum chemistry are shown in table 1 and 2 respectively. There were no significant difference ($P > 0.05$) amongst the WBC values of the 6 groups and the respective values are within the normal range for rabbits indicating the absence of negative effect of the diets on the experimental animal. However, there were significant differences ($P < 0.05$) between the other parameters with RBC ranging from 4.2 – 4.8/mm³ and T3 showing significant difference from the other groups. There were also significant differences ($P < 0.05$) in the packed cell volume (PCV) and the MCV among the treatments.

The results further showed significant differences ($P < 0.05$) in the Mean Corpuscular Haemoglobin (MCH). In table 2, there were no statistically significant differences amongst the various feed treatments ($p > 0.05$) on the protein concentration, cholesterol levels, ALT, AST and potassium levels. However, apart from T1 and T2, there were significant differences ($P < 0.05$) on the glucose levels between the various treatments.

The results of this study is in agreement with Ogungbesan *et al.* (2014) who monitored blood profiles as influenced by *Gliricidia* leaf meal (GLM) supplemented with Maxigrain® enzyme in laying hens also concluded that supplementation of feed with Maxigrain® enzyme caused no discomfort or

diseases to the animals, furthermore, and most importantly the combination is of high health importance because of low cholesterol content. Blood is known to be vital to the life of an organism. This is a medium through which nutrients are conveyed to various parts of the body system of an organism. A readily available and fast means of assessing clinical and nutritional status of an animal on feeding trial may be the use of blood analysis (Olabanji *et al.*, 2007) [12]. Haematological

parameter is an important and reliable medium used to monitor and evaluate health and nutritional status of animals (Babatunde *et al.*, 1992; Onifade, 1993; Gupta, *et al.*, 2007) [3, 14, 6]. Therefore, the absence of any negative effects of the experimental diets used in this study suggested that the experimental diets were able to supply the required nutrition which maintained the normal blood parameters of the experimental animals.

Table 2: Effects of experimental diets on hematological parameters of growing rabbits

Parameters	Inclusion levels of Dried Rumen Digesta						
	(0%)		(12.5%)		(25%)		
	T1 (M)	T2 (M+E)	T3 (12.5%RD)	T4 (12.5%RD+E)	T5 (25%RD)	T6 (25%RD+E)	
RBC/mm ³	4.2±1.1 ^a	4.6±0.6 ^a	4.8±0.8 ^a	3.9±1.2 ^b	4.6±0.6 ^a	4.5±1.3 ^a	**
WBC/L	7.7±1.1	8.1±2.4	8.2±1.1	7.5±0.	8.0±1.1	8.4±0.9	NS
PCV (%)	42±1.5 ^a	40±1.2 ^a	38±0.9 ^b	40±1.2 ^a	36±1.5 ^b	39±1.0 ^a	**
MCV (mm ³)1.2 ^b	91.30±0.8 ^a	83.33±	97.44±0.9 ^a	86.96±1.6 ^b	94.43±1.2 ^a	95.12±0.8 ^c	**
Hb (g/dl)	12.8±2.0	11.3±0.	10.6±1.4	12.2±2.2	11.6±0.6	2.0±1.8	NS
MCH (pg/cell)	27.83±1.4 ^a	23.54±1.6 ^b	27.18±1.2 ^a	26.53±2.1 ^a	28.52±1.7 ^a	26.27±2.1 ^a	NS
MCHC (%) 1.6 ^a	30.48±0.4 ^a	28.25±	27.89±0.8 ^b	30.5±1.1 ^a	32.22±1.6 ^a	30.77±1.2 ^a	NS

^{abc} means within the same rows bearing different superscripts are significantly different ($p<0.05$).

T1 control (0% DRD without enzyme) T2 control (0% DRD with enzyme), T3 12.5% DRD without enzyme, T4 12.5% DRD with enzyme, T5 25% DRD without enzyme, T6 25% DRD with enzyme, ± SEM, (NS) not significantly different among the six(6) treatment, (**) there is significant different among the six(6) treatment.

Table 3: Effects of the experimental diets on serum chemistry parameters of growing rabbits

Parameters	Inclusion levels of Dried Rumen Digesta						
	(0%)		(12.5%)		(25%)		
	T1 (M)	T2 (M+E)	T3 (12.5%RD)	T4 (12.5%RD+E)	T5 (25%RD)	T6 (25%RD+E)	
Prot(g/dl)	5.2±0.24	5.3±0.42	5.1±0.82	5.36±1.0	5.35±0.6	5.45±0.28	NS
Glu(mg/dl)	145±2.2 ^a	142±2.8 ^a	133±1.2 ^b	136±2.6 ^b	130±1.8 ^c	130±0.8 ^c	**
Chol. (mg/dl)	32.2±1.6	30.8±2.4	31.8±1.6	33.6±1.9	38.5±2.12	40.4±1.65	NS
ALT (IU/L)	60±1.6	64±2.2	62±0.8	66±0.6	65±0.8	64±0.4	NS
AST (IU/L)	68±1.31	67±0.9	66±1.18	66±1.34	68±0.8	64±0.9	NS
ALP (IU/L)	42±0.2 ^a	44±0.6 ^a	48±0.12 ^a	42±0.84 ^a	75±0.18 ^b	79±0.9 ^b	**
Na ⁺ (mmol/L)	130±2.32 ^a	138±1.61 ^a	128±0.91 ^a	133±1.42 ^a	98±0.6 ^b	92±1.0 ^b	**
K ⁺ (mmol/L)	3.1±0.2	3.3±0.2	3.3±0.2	3.6±0.42	3.5±0.3	3.4±0.2	**

^{abc} means within the same column bearing different superscripts are significantly different ($p<0.05$).

T1 control (0% DRD without enzyme) T2 control (0% DRD with enzyme), T3 12.5% DRD without enzyme, T4 12.5% DRD with enzyme, T5 25% DRD without enzyme, T6 25% DRD with enzyme, ± SEM, (NS) not significantly different among the six(6) treatment, (**) there is significant different among the six(6) treatment.

Conclusion

This study showed that from the various parameters monitored, rumen digesta can be used with or without enzyme supplementation as partial replacement for maize in providing energy and protein requirements for weaner rabbits without any significant effects on hematological and serum chemistry parameters. It can be concluded that rumen digesta can be included up to 25% with or without enzyme supplementation in maize concentrate diets for rabbits without any adverse effects. However, further investigation on carcass quality and histopathology of some organs are necessary as these will help to provide further information on the health status of the rabbits.

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