

Growth and reproductive performance of rabbit bucks fed replacement levels of fermented *Jatropha* (*Jatropha carcass*) seed meal

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Abstract

*This study was conducted to investigate the effect of fermented *Jatropha* seed meal on growth and reproductive characteristics of rabbit bucks. Thirty (30) (twelve weeks old) cross bred rabbits (American chinchilla × New Zealand white) were used for the experiment. The rabbits were randomly assigned to five dietary treatments replicated three times with two rabbits per replicate in a completely randomised design (CRD). The five experimental diets were compounded using fermented *Jatropha* seed meal (FJSM) to replace soya bean meal at 0, 25, 50, 75 and 100%, respectively. The experiment lasted for 12 weeks. Data collected during the feeding trial include final body weight, total weight gain, total feed intake, feed conversion ratio and mortality. At the end of the feeding trial, 3 bucks per treatment were slaughtered and the reproductive organs dissected out for testicular and epididymal morphometrics as well as gonadal sperm assessment. All the data collected during the experiment were subjected to analysis of variance. The result of the growth performance showed that final body weights and total weight gain were highly ($P < 0.001$) influenced by the replacement levels of fermented *Jatropha* seed meal. Final body weight of buck decreased as the level FJSM increased in the diets. Among the fermented *Jatropha* seed meal-based diets, rabbit fed T2 (25% FJSM) recorded significant higher ($P < 0.05$) final body weight (2011.10g) than the other group. The results of epididymal and testicular characteristics as well as testicular sperm characteristics were significantly ($P < 0.001$) influenced by replacement levels of FJSM in the diets. It was concluded from this study that feeding rabbit buck with replacement levels of fermented *Jatropha* seed meal beyond 25% adversely affected growth and reproductive performance. Therefore, it is recommended that fermented *Jatropha* seed meal at 25% replacement levels can be used in rabbit diets.*

keywords: Buck, Fermentation, Growth, *Jatropha*, Rabbits, Reproductive Performance

La Croissance et La performance reproductrice des mâles lapins nourris avec des niveaux de remplacement de farine de graines de *jatropha* fermenté (carcasse de *Jatropha*)



Résumé

*Cette étude a été menée pour étudier l'effet de la farine fermentée de graines de *Jatropha* sur la croissance et les caractéristiques reproductrices des mâles lapins. Trente (30) (douze semaines) lapins croisés (chinchilla américain × Nouvelle-Zélande) ont été utilisés pour l'expérience. Les lapins ont été assignés au hasard à cinq traitements diététiques répliqués trois fois avec deux lapins par réplique dans une conception complètement randomisée (le 'CRD'). Les cinq régimes expérimentaux ont été composés utilisant le repas fermenté de graine de *Jatropha* (le 'FJSM') pour remplacer le repas de haricot de soja à 0, 25, 50, 75 et 100%, respectivement. L'expérience a duré 12 semaines. Les données recueillies au cours de*

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*l'essai d'alimentation comprennent le poids corporel final, le gain de poids total, la consommation totale d'aliments pour animaux, le ratio de conversion des aliments pour animaux et la mortalité. À la fin de l'essai d'alimentation, 3 dollars par traitement ont été abattus et les organes reproducteurs disséqués pour l'évaluation testiculaire et épидидymalmorphométrique ainsi que l'évaluation du sperme gonadal. Toutes les données recueillies au cours de l'expérience ont fait l'objet d'une analyse de la variance. Le résultat de la performance de croissance a prouvé que les poids corporels finaux et le gain total de poids ont été fortement ($P < 0.001$) influencés par les niveaux de remplacement du repas fermenté de graine de *Jatropha*. Le poids corporel final de buck a diminué pendant que le niveau FJSM augmentait dans les régimes. Parmi les régimes fermentés à base de farine de graines de *Jatropha*, le T2 nourri au lapin (25 % de 'FJSM') a enregistré un poids corporel final ($P < 0,05$) plus élevé (2011,10 g) que l'autre groupe. Les résultats des caractéristiques épидидymales et testiculaires aussi bien que des caractéristiques testiculaires de sperme ont été sensiblement ($P < 0.001$) influencés par des niveaux de remplacement de FJSM dans les régimes. Il a été conclu de cette étude que l'alimentation de l'argent du lapin avec des niveaux de remplacement de farine fermentée de graines de *Jatropha* au-delà de 25% a nui à la croissance et la performance reproductive. Par conséquent, il est recommandé que le repas fermenté de graine de *Jatropha* aux niveaux de remplacement de 25% puisse être employé dans des régimes de lapin.*

Mots-clés: mâles lapins, fermentation, Croissance, *Jatropha*, Performance reproductive

Introduction

The scarcity of conventional feedstuff most especially protein and energy sources in most developing countries of the world has continued to be one of the major challenges facing livestock producers (Biobaku and Dosunmu, 2003). Searching for alternative feedstuffs is paramount if animal production and supply of quality and quantity animal protein is to be sustained (Aro *et al.*, 2009). Rabbit production has been identified as one of the fastest and most efficient means of attaining animal protein self-sufficiency (Yakubu and Wafar, 2014). This is attributed largely to the rabbit's high rate of reproduction, early maturity, rapid growth rate, efficient converter of feed to meat of high nutritional value (Ghosh *et al.* 2008). The production efficiency of commercial rabbit farms is largely dependent on the quality of semen the buck produces, litter size at kindling and the survivability of the bunnies up to weaning age (Odeyinka *et al.* 2008). In addition, the pre-weaning growth is very critical in meat rabbits due to its impact on

the meat produced at the finisher stage of production (Gerencser *et al.*, 2011). As production directly depends on reproduction, the reproductive performance of rabbits becomes an important aspect in determining the profitability of commercial rabbit breeding (Lazzaroni *et al.* 2012). However, nutrition and high cost of feed are among other factors affecting rabbit production. High cost of feed has been identified as an impending factor affecting successful rabbit production in most developing countries of the world (Apori *et al.*, 2014). The search for alternative plant protein sources over the years as a replacement for conventional feedstuffs has been the focus of animal nutritionist, physiologist and breeders (Owen *et al.*, 2010; Wafar and Tarimbuka, 2016). The use of non-conventional protein sources such as pigeon pea (Ahamefule 2005), mucuna seed meal (Carew *et al.*, 2003), cotton seed meal (Amao and Showunmi 2016), Kapok seed meal (Wafar *et al.*, 2017) in animal ration is well documented. One of the non-conventional feedstuffs focused in this

study is *Jatropha curcas* seed. *Jatrophas* are oil plants belonging to the *Euphorbiaceae* family. It has both toxic and non-toxic genotypes while *Jatropha platyphylla* is a non-toxic species (Makkar *et al.*, 2012). It is known among some Nigerian ethnic groups as *Bindazugu* (Hausa), *Kolkolaje*, (Fulfulde), *Lapalapa* (Yoruba), *Wuluidu* (Igbo) and *Safudruna* (Higgi). They are drought-resistant perennial and multipurpose shrubs, similar to the cassava plant (Elbehri *et al.*, 2013). The seed has been reported to contain 56.4% crude protein higher than that of full fat soya bean meal (48%) (Makkar and Becker, 1999). This relatively high protein content of *Jatropha curcas* can be advantageous since it is not utilised by human beings as source of food (Hammarneh *et al.*, 2012). However, *Jatropha* seed meal also contains anti-nutritional factors; such as lectin, trypsin inhibitor, saponin, phytate, and phorbol esters (Makkar and Becker, 1998, Antevy *et al.*, 2017). Phorbol ester is considered as the most toxic anti-nutritional factor. The use of *Jatropha* in animal nutrition is however limited due to these anti-nutrients (Makkar and Becker, 1999). The use of processing methods such as cooking, toasting, fermentation and soaking have been reported to reduce the contents of anti-nutritional factor in *Jatropha curcas* seed meal (Antevy *et al.*, 2017). A study conducted by Antevy *et al.* (2017) to evaluate the performance of broiler chickens fed differently processed *Jatropha curcas* seed meal showed that broiler chicken fed fermented *Jatropha* seed meal recorded higher feed intake, final body weight gain and superior feed conversion ratio. Fermentation is one of the oldest forms of feed processing and preservation Ross *et al.* (2002), and also has the capacity to improve nutritional and functional properties of the feed stuff such as improving the digestibility, flavor and

reduce the levels of anti-nutrients in tropical seed meals thereby increasing the possibilities of its utilization. (Hotz and Gibson, 2007 and Frias *et al.* 2008) However, there is little or no information on the use of fermented *Jatropha* seed meal on the growth and reproductive parameter of rabbit buck. This study therefore was designed to assess the effects of fermented *Jatropha* seed meal on growth and reproductive performance of rabbit bucks.

Materials and methods

Location of the study

This study was carried out at the Rabbit Unit of the Department of Animal Science and Range Management, Modibbo Adama University of Technology, Yola Adamawa State. Yola is between latitude 7° 11' North and Longitude 11° 14' East and at an elevation of 364m above sea level in the north eastern part of Nigeria. It has mean relative humidity ranges from 30 - 50% with a minimum in February to March and as low as 10% and a maximum of about 90% in August. The maximum temperature can reach 38°C particularly in April, while minimum temperature can be as low as 18°C (Adebayo, 1999).

Collection and processing of *Jatropha curcas* seeds

Jatropha curcas fruits were purchased from Yola market. The fruits were cracked mechanically to remove the seeds and processed according to modified method of Wafar *et al.* (2019). Five kg of the seeds was packaged in a jute bag and placed in a pot containing 20 litres of water then cooked for 45 minutes. The use of jute bag is to control leaching and possibility of hydrolysis of some nitrate. The pot was placed on a tripod stand having firewood as a source of heat. The cooked seeds were packed in a polythene bag to allow natural fermentation to take place. The fermentation process lasted for 120 hours (5 days). The fermented *Jatropha* seeds were

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sundried and milled using hammer milling machine to produce fermented *Jatropha* seed meal (FJSM).

Experimental animals, management, design and diets

Thirty buck rabbits (American chinchilla × New Zealand white) with an average weight of (1140±2.0g) were procured within Yola metropolis. They were assigned to five dietary treatments. Replicated three times with two rabbits per replicate in a completely randomised design (CRD). Each rabbit was housed in a cage measuring 150cm x100cm x120cm fitted with aluminium feeder and drinker. The experimental animals were given

prophylactic treatment against endo and ecto-parasites using 0.3ml/kg Ivomec®.

Five diets were compounded using fermented *Jatropha* seed meal (FJSM) to replace soya bean meal at 0, 25, 50, 75 and 100% designated as treatment 1, 2, 3, 4 and 5 respectively as shown in Table 1. The experiment lasted for 12 weeks.

Laboratory analysis

Proximate composition and anti-nutritional composition of raw and fermented *Jatropha* seed meal were determined (dry matter (DM), crude protein (CP), crude fibre (CF), ether extracts (EE) and Ash content) as described by AOAC (2010). The extraction of phorbol ester was carried out using modified method of Makkar *et al.* (1997).

Table 1: Ingredient and percentage composition of the experimental diets

Ingredients	Replacement levels of fermented <i>Jatropha</i> seed meal				
	T1 (0 %)	T2 (25%)	T3 (50 %)	T4 (75%)	T5 (100 %)
Maize	52.00	52.00	52.00	52.00	52.00
SBM*	18.00	13.50	9.00	4.50	0.00
FJSM**	0.00	4.50	9.00	13.50	18.00
Groundnut haulms	10.00	10.00	10.00	10.00	10.00
Maize offal	13.00	13.00	13.00	13.00	13.00
Fish meal	3.00	3.00	3.00	3.00	3.00
Bone meal	3.00	3.00	3.00	3.00	3.00
Premix*	0.25	0.25	0.25	0.25	0.25
Lysine	0.25	0.25	0.25	0.25	0.25
Salt	0.25	0.25	0.25	0.25	0.25
Methionine	0.25	0.25	0.25	0.25	0.25
Total	100	100	100	100	100
Calculated analysis					
Crude protein	16.83	16.64	16.73	16.65	16.70
Crude fibre	8.84	8.76	8.70	8.63	8.69
Ca	1.23	1.22	1.20	1.23	1.24
P	0.90	0.90	0.91	0.89	0.83
Lysine	0.91	1.89	0.88	0.97	0.89
Methionine	0.58	0.63	0.61	0.59	0.60
***ME Kcal/kg	2978.14	2998.13	2990.14	2992.88	2991.34

*Vitamin-mineral premix provided the following per kg of feed: Vit. A 1500 IU; Vit.D₃ 3000 IU; Vit.E 30 IU; Vit.K 2.5mg; Thiamine B₁ 3mg; Riboflavin B₂ 6 mg; Pyridoxine B₆ 4 mg; Niacin 40 mg; Vit. B₁₂ 0.0 mg; Pantothenic acid 10mg; Folic acid 1mg; Biotin 0.08 mg; Chloride 0.125mg; Mn 0.0956 g; Antioxidant 0.125 g; Fe 0.024 g; Cu 0.006 g; Se 0.24 g; Co 0.24.

SBM *= Soya bean meal, FJSM** = Fermented *Jatropha* seed meal, ME**= Metabolizable energy

Data collection

Growth parameter

The growth parameter determined were feed intake, weight gain and FCR was

calculated as ratio of total feed intake to total weight gain. Feed intake was calculated as the difference between feed offered and left over after a period of 24hours

Evaluation of reproductive parameter

At the end of 12 weeks of feeding trial, 3 bucks per treatment were randomly selected and slaughtered for reproductive organs evaluation.

Testicular morphometry

After slaughtering, the epididymis was trimmed off the testis; the right and left testis were weighed using a digital scale (MLP 231 scale). The length and width of testis were measured using a pair of vernier caliper while the volume of the testis was determined using water displacement method of Archimede's principle using a measuring cylinder.

The testicular morphometry parameter evaluated were:

Paired testis weight (g), left testis weight (g), right testis weight (g), mean testis length (cm)

Left testis length (cm), right testis length (cm), left testis width (cm), right testis width (cm). Paired testis volume (ml), left testis volume (ml), right testis volume (ml).

Epididymal morphometry

Epididymal parameters include: left epididymal weight (g), right epididymal weight (g), left epididymal length (cm), right epididymal length (cm)

Sperm analysis

Sperm morphology was determined according to the method described by Zemjanis (1970). A smear of the semen was made by cutting the right testis along the equatorial region and rubbing the cut surface on a clean glass slide. Two drops of eosin-nigrosin dye that had been thoroughly mixed was added. Another smear was made on a glass slide and viewed under a light microscope to identify normal and abnormal cells on the slide. The normal cells were expressed as the percentage of number of cells counted on each field of the slide (Ogbuewu, 2008). Sperm count was determined haemocytometrically by homogenization technique as described by Adejumo (2006) and Amao *et al.* (2012).

The tunica albuginea was carefully removed from the testis. A portion of the parenchyma tissue was taken and homogenized by maceration with a pair of sharp scissors for 5 minutes in a beaker containing 10 ml of physiological saline solution. The homogenate was filtered through a double layer cheese cloth and the filtrate diluted to ratio 1:20 with de-ionized water. Some drops of the homogenate were introduced into an improved Neubauer haemocytometer counting chamber. All the elongated spermatids and mature sperm cells in the four diagonal and the centre squares of the haemocytometer were counted in each diluted homogenate.

Motility was determined by a modification of the method of Ogbuewu *et al.* (2008). A drop of the homogenate was placed on a sterile slide, covered with a cover slip and observed under the microscope at X 400 Magnification and scored between 0 and 100%.

The parameters evaluated were sperm count, motile and non- motile sperm percentages, sperm volume and concentration, normal and abnormal sperm, round and elongated spermatids.

Statistical analysis

All data obtained were subjected to analysis of variance (ANOVA) using JMP SAS (2013) software where significant differences existed, Duncan's Multiple Range Test (DMRT) option of same software were used to separate means.

Results and discussion

Proximate and antinutritional composition of raw and fermented Jatropha Seed meal

The proximate and anti-nutritional compositions of the raw and fermented Jatropha seed meal (FJSM) are presented in Table 2. The results showed FJSM recorded higher values of crude Protein (CP) 43.67%, ash (9.13%) and lower values of crude fibre (2.41%), ether extracts

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(20.18%), nitrogen free extracts (NFE) (14.52%) and metabolizable energy (3765.83%), respectively. The crude fibre (CF) and ether extracts (EE) values ranged from 10.30 – 14.32% and 2.17- 4.76%, /respectively. Nitrogen free extract for raw and fermented jatropha meals were 48.61 and 48.14% respectively. The result of the anti-nutritional factors (ANFs) composition however, shows that raw Jatropha seed contained 3.32mg/100g phorbolster, 20.45mg/100g trypsin inhibitors, 191.56mg/100g saponnin, 278.67mg/100g tannin, 94.67mg/100g oxalates and 234.56mg /100g phytates, while the fermented Jatropha seed recorded 2.11mg phorbolster, 8.67mg/100g trypsin inhibitors, 78.96mg/100g saponnin, 87.89mg/100g tannins 56.45mg/100g oxalates and 58.11mg/100g phytates. The results showed decrease in antinutrients as a result of fermentation. The higher crude protein (CP) content observed in fermented Jatropha seed meal could be attributed to the effect of the fermentation process that

leads to crude protein improvement. The CP values of both raw and fermented Jatropha seed meal were within the range of 35-50% crude protein reported by Aslani *et al.* (2007). This indicates that fermented Jatropha seed meal may be comparable to soybean meal in its usefulness in rabbit diets (Makkar *et al.*, 1998). Presences of anti-nutrients in the raw jatropha seed meal confirm the earlier report of Makkar and Becker, (1999) that jatropha seed meal contained high concentration of anti-nutritional factors (ANFs). The use of fermentation as a method of detoxification however, reduced the concentration of ANFs and increased the nutrient composition of Jatropha seed meal, but did not completely remove the ANFs in the seed. The finding is in line with earlier reports of Makkar *et al.* (1998), Wafar (2013) and Yakubu *et al.* (2017) when they subjected tropical legume seeds to different processing methods and recorded partial reduction of ANFs in the seeds.

Table 2: Proximate composition of raw and fermented Jatropha seed meal

Constituents (%)	Raw	Fermented	
Dry matter	91.11	89.91	
Crude Protein	30.11	43.67	
Ether extracts	32.13	20.18	
Crude fibre	4.06	2.41	
Ash	7.13	9.13	
NFE	17.68	14.52	
*Metabolizable Energy	4344.24	3765.83	
<i>Antinutritional factor (mg/100g)</i>			<i>% Reduction</i>
Phorbolsters	3.32	2.11	63.55
Trypsin inhibitors	20.45	8.67	42.39
Saponin	191.56	78.98	41.22
Tannins	278.67	87.89	31.53
Oxalates	94.67	56.45	59.62
Phytates	234.56	58.11	24.77

*Metabolizable Energy = ME (kcal/kg) = 37 x % CP + 81 x % EE + 35.5 x % NFE. Calculated according to the formula of Pauzenga, (1985)

CP= Crude protein EE = Ether extracts NFE = Nitrogen free extracts

Growth performance of buck rabbits fed replacement levels of fermented Jatropha Seed meal

Table 3, showed the growth performance of

buck rabbits fed replacement levels fermented Jatropha seed meal. The final body weights and total weight gain were highly (P<0.001) influenced by the

replacement levels of fermented *Jatropha* seed meal. Final body weight of buck decreased as the level FJSM increased in the diets. Among the fermented *Jatropha* seed meal based diets rabbit fed T2 (25% FJSM) recorded significantly higher ($P<0.05$) final body weight 2011.10g. Total body weight gain and total feed intake varied from 217.07g in T5 (100% FJSM) to 1330.55g in T1 (0% FJSM) and 1206.51g in T1 (25% FJSM) to 4325.70g in T1 (0%FJSM) respectively. Feed conversion ratio also differed significantly ($P<0.05$) across the treatments. It was observed that the superior feed conversion was recorded T1 (0%FJSM) 3.25 while those fed T5 (100%FJSM) recorded poor feed conversion ratio. Rabbit fed T5 had higher mortality (3.00%). It was observed from the study that rabbits' bucks fed T3 (50% FJSM), T4 (75% FJMS) and T5 (100% FJSM) recorded lower feed intake. The low feed intake could be attributed to the residual effect of anti-nutritional factors in the fermented *Jatropha* seed meal which resulted to low palatability of diets. Tannins and saponins have reported to decrease feed intake as a result of its astringent properties (Ogbu *et al.* (2015). Weaver and Kanna,

(2002) in their study also reported that phytate reduce the bioavailability of divalent cations due to insoluble complexes formation during digestion and absorption of minerals. This implied that rabbit buck fed diets containing 50, 75 and 100% FJSM replacement levels could not efficiently absorb the dietary nutrients available in the gastro-intestinal tract as a result of high concentration of these residual antinutritional factors. Esonu *et al.* (2001) reported that high trypsin inhibitors in the diet reduce protein digestibility resulting in poor utilization of available nutrients. Phorbol esters are toxic to livestock even at very low concentrations (Goel *et al.*, 2007). The result from this study confirmed earlier findings of Abdel-Shafy *et al.* (2011) who reported significant reduction in feed intake and growth rate of more than 30% as a result of residual phorbol esters in *Jatropha* seed meal. Rabbit fed T5 (100% FJSM) recorded high percentage mortality this could be attributed to residual content of phorbol ester. Agboola and Adenuga, (2015) reported high percentage of mortality of birds fed 10% inclusion level of *Jatropha* seed meal

Table 3: Growth Performance of Buck rabbits Fed Replacement levels of Fermented *Jatropha* Seed meal

Parameter	Replacement levels of fermented <i>Jatropha</i> seed meal					SEM	P- value
	T1 (0%)	T2 (25%)	T3 (50%)	T4 (75%)	T5 (100%)		
Initial body weight (g)	1225.29	1195.34	1205.53	1140.47	1188.41	21.73	0.12
Final body weight (g)	2555.83 ^a	2011.10 ^b	1554.85 ^c	1623.57 ^c	1405.48 ^d	34.34	0.01
Total weight gain (g)	1330.55 ^a	815.76 ^b	344.32 ^{cd}	484.11 ^c	217.07 ^d	30.86	0.02
Total feed intake (g)	4325.70 ^a	3599.06 ^b	1531.02 ^d	2565.67 ^c	1206.51 ^e	47.00	0.01
Feed conversion ratio	3.25 ^b	4.41 ^{ab}	4.57 ^{ab}	5.30 ^{ab}	5.58 ^a	0.38	0.04
Mortality (%)	0	1.00	1.00	1.00	3.00	-	-

Means in the same row bearing different superscripts differ significantly

SEM = Standard error mean, ns = not significant

Testicular characteristics of rabbit bucks fed fermented levels of fermented seed meal

The result showed significant variations by the levels replacement (Table 4). The values recorded for the right testis and left testis

weights were within the range of 1.00 – 1.80g, 1.05 g – 1.84g and 2.05 – 3.64g respectively. Bucks fed *Jatropha* based diets especially beyond 25% inclusion had significant lower weight testis weights. From the result it appeared the inclusion of

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fermented *Jatropha* seed meal in the diet relatively decreased the weight of the testis. The ability of the testes to store spermatozoa is of immense importance in rabbit breeding programme. The decrease in testicular sperm reserve observed on rabbit fed T3 (50%), T4 (75%) and T5 (100%) fermented *Jatropha* seed meal is a pointer that higher inclusion of the seed meal negatively affected testicular sperm reserve. This indicates that feeding fermented *Jatropha* seed meal in rabbit diet beyond 25% is detrimental to the development of spermatogenic potentials of the buck as it has been observed in this present study. Testis size is a good indicator of the present and future spermatozoa production of an animal (Perry and

Petterson, 2001; Gupta and Mohanty, 2003; Togun and Egbunike, 2006). The knowledge of basic morphometric characteristics of the reproductive organs have been found to provide valuable information in the evaluation of breeding and fertility potential of the animals (Galmessa *et al.*, 2003). Larger testes (without any abnormality) have been reported to produce more spermatozoa than smaller testes (Oyeyemi *et al.*, 2002;). Morton (2006) reported that in sacrificed animals, decreased weight of the testes indicates widespread or diffuse loss of seminiferous epithelial cells. The testes which possess greater number of sertoli cells were heavier and produced more spermatozoa than testes with fewer sertoli cells (Britto *et al.*, 2004).

Table 4: Testicular characteristics of rabbit buck fed replacement levels of fermented seed meal

Parameter	Replacement levels of fermented <i>Jatropha</i> seed meal					SEM	P-value
	T1 (0%)	T2 (25%)	T3 (50%)	T4 (75%)	T5 (100%)		
Right testis weight (g)	1.80 ^a	1.51 ^b	1.04 ^b	1.03 ^c	1.00 ^d	0.38	0.05
Left testis weight (g)	1.84 ^a	1.43 ^b	1.30 ^b	1.50 ^b	1.05 ^b	0.37	0.03
Paired testis weight (g)	3.64 ^a	2.94 ^b	2.34 ^c	2.53 ^c	2.05 ^c	0.38	0.04
Left testis length (cm)	2.45 ^a	2.16 ^b	1.23 ^c	1.15 ^c	1.03 ^c	0.92	0.05
Right testis length (cm)	2.40 ^a	2.01 ^a	1.13 ^b	1.02 ^b	1.02 ^b	0.44	0.03
Mean testis length (cm)	2.43 ^a	2.08 ^b	1.18 ^c	1.08 ^c	1.02 ^c	0.71	0.05
Left testis width (cm)	1.93 ^a	1.79 ^a	1.13 ^b	1.09 ^b	1.02 ^c	0.30	0.04
Right testis width (cm)	1.79 ^a	1.68 ^b	1.09 ^c	1.07 ^c	1.05 ^c	0.29	0.02
Mean testis width (cm)	1.86 ^a	1.74 ^b	1.11 ^c	1.08 ^c	1.03 ^c	0.49	0.05
Right testis volume (ml)	3.19 ^a	3.15 ^a	2.87 ^b	2.79 ^b	2.18 ^b	0.65	0.04
Left testis volume (ml)	3.30 ^a	3.40 ^a	2.74 ^b	2.80 ^b	2.82 ^b	0.64	0.03
Paired testis volume(ml)	3.25 ^a	3.28 ^a	2.80 ^b	2.79 ^b	2.50 ^b	1.48	0.02

Means in the same row bearing different superscripts differ significantly
SEM = Standard error mean

Epididymal characteristics of rabbit bucks fed graded levels of fermented seed meal

Table 5, shows the epididymal characteristics of rabbit bucks fed graded levels of fermented seed meal. The result revealed all the parameters evaluated were significantly influenced. Mean epididymis weight and mean epididymis length were significantly affected by replacement levels of fermented *Jatropha* carcass seed meal in the diets. The mean epididymis weight of the bucks on the T1 (0% FJSM) 2.73g and

T2 (25% FJSM) 2.74g were significantly ($p < 0.05$) heavier compare to those on T3 (50% FJSM) 1.79 g, T4 (75% FJSM) 1.74 g and T5 (100% FJSM) 1.60 g. The right and left and epididymis weights of rabbit bucks fed replacement levels of fermented *Jatropha* seed meal are within the range of 1.74 – 2.74g, 1.46 – 2.75g and 1.60-2.73g respectively. The higher epididymis weight of rabbits on T1 (0.0% FJSM) and T2 (25% FJSM) implies that the bucks have the ability to store spermatozoa. Smaller

epididymis weights of the bucks on T3 (50%), T4 (75% FJSM) and T5 (100% FJSM) inclusion levels of fermented *Jatropha* seed meal suggest fewer spermatozoa is being stored (Perry and Petterson, 2001). The higher epididymis weight of observed in rabbits fed T1 (0% FJSM) and T2 (25% FJSM) were within the normal weight of epididymis (2- 3.0g) reported by Olomu *et al.*, (2019) for buck rabbits raised under tropical condition. This

implies that fermented *Jatropha* seed meal promotes the growth and development of testicular glands and gonadal sperm reserve. Colenbrander and Kemp (1990) reported that testicular weight and quantity of sperm produced are correlated. Morris *et al.* (1999) also stated that within a species of animals, there often exist positive correlation between spermatozoa production, testicular size and testicular length.

Table 5: Epididymal characteristics of rabbit buck fed Replacement levels of fermented seed meal

Parameter	Replacement levels of fermented <i>Jatropha</i> seed meal					SEM	P-value
	T1 (0%)	T2 (25%)	T3 (50%)	T4 (75%)	T5 (100%)		
Left epididymal weight (g)	2.75 ^a	2.73 ^a	1.85 ^b	1.74 ^b	1.46 ^c	0.68	0.04
Right epididymal weight (g)	2.70 ^a	2.74 ^a	1.73 ^b	1.75 ^b	1.74 ^b	0.64	0.02
Mean epididymal weight (g)	2.73 ^a	2.74 ^a	1.79 ^b	1.74 ^b	1.60 ^b	1.09	0.05
Left epididymal length (cm)	17.35 ^a	16.30 ^b	14.29 ^c	13.31 ^d	11.27 ^e	0.69	0.03
Right epididymal length (cm)	17.05 ^a	15.89 ^a	14.10 ^b	13.29 ^c	11.18 ^d	0.61	0.02
Mean epididymal length (cm)	17.20 ^a	16.09 ^a	14.19 ^b	13.30 ^b	11.22 ^c	1.10	0.02

Means in the same row bearing different superscripts differ significantly SEM = Standard error mean

Testicular sperm characteristics of rabbit buck fed replacement fermented levels of *Jatropha* seed meal

The results of testicular sperm characteristics are presented in Table 6. The result showed significant variation (P<0.001) across the dietary treatments. Rabbit T1 (0% FJSM) 183.12×10^6 recorded significant higher (P<0.001) sperm cell count but similar to those on T2 (25% FJSM) 170.67×10^6 compare to those fed T3 (50% FJSM), T4 (75% FJSM) and T5 (100% FJSM) diets. The values for motile and non-motile sperm were between 30.12 in T5 (100%) – 76.22% in T1 (0.00%) and 10.45 in T1 (0%) to 57.67% in T5 (100%). Buck fed T1 (0%) fermented *Jatropha* seed meal recorded the least non motile value of 10.45%. The result showed that motile sperm decrease progressively as the level of fermented *Jatropha* seed meal increased in the diet, while abnormal sperm percentages increase progressively with increase in levels of fermented *Jatropha* seed meal. One of the methods of assessing

reproductive efficiency of the buck rabbit is through the measurement of semen quality (Ogbuewu *et al.*, 2008). Every animal species has its capacity for sperm production, which is determined genetically but it has however, been clearly observed that other factors like nutrition, disease and stress influence the portion of the germinal epithelium that enters into spermatogenesis (Harper *et al.*, 1999). The results of this study clearly indicate that dietary inclusion of fermented *Jatropha* seed meal beyond 25% negatively affect the testicular sperm characteristics of buck rabbits. Nutrition has long ago been established in affecting the secretory functions of the accessory sex glands, the products of which constitute the seminal plasma (Oyeyemi and Okediran, 2007). The progressive decrease of sperm cell count on rabbit fed graded levels of fermented *Jatropha* seed meal agreed with the findings of Ogbuewu *et al.* (2008) that secretory functions of the accessory sex glands are very sensitive to dietary changes and that

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the slight changes in feed chemical composition goes a long way in influencing sperm quantity. Sperm motility is an important index in reproductive assessment because it demonstrates the ability of sperm to move and fertilize an ovum (Ogbuewu, 2008). Significant higher number of motile

sperm cells in buck rabbits fed T1 (0.0% FJSM) and T2 (25% FJSM) could be attributed to sperm cell development in the seminiferous tubules, in the sense that spermatozoa may need a moderate but progressive development for them to have excellent motility.

Table 6: Testicular sperm characteristics of buck rabbit buck fed replacement levels of fermented *Jatropha* seed meal

Parameters	Replacement levels of fermented <i>Jatropha</i> seed meal					SEM	P-value
	T1(0%)	T2 (25%)	T3 (50%)	T4 (75%)	T5 (100%)		
Sperm count (x10 ⁶)	183.12 ^a	170.67 ^{ab}	130.11 ^b	120.00 ^c	100.67 ^d	5.78	0.02
Motile sperm (%)	76.22 ^a	61.67 ^b	40.11 ^c	35.23 ^c	30.12 ^c	1.01	0.04
Non motile sperm (%)	10.45 ^d	20.45 ^c	30.13 ^b	56.45 ^a	57.67 ^a	2.52	0.01
Normal sperm (%)	70.11 ^a	64.23 ^b	20.34 ^c	20.11 ^c	18.67 ^c	1.20	0.03
Abnormal sperm (%)	10.56 ^c	21.56 ^c	32.54 ^b	34.11 ^b	40.11 ^a	4.81	0.03

Means in the same row bearing different superscripts differ significantly

SEM = Standard error mean

Conclusion and recommendation

It was concluded from this study that feeding rabbit buck with replacement of fermented *Jatropha* seed meal beyond 25% adversely affected growth and reproductive performance. Therefore, it is recommended that fermented *Jatropha* seed meal at 25% replacement levels can be used in rabbit diets.

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