Effects of substitution of groundnut meal by defatted and boiled *Jatropha curcas* seed kernel in diet of broiler chicks in Senegal

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Abstract

This study was conducted to determine, in a growth diet, the effect of a substitution of groundnut meal by *Jatropha curcas* kernel meal physico-chemically treated on viability, feed intake and growth performances of broiler chicks.

The feed experiment lasted for seven days with twenty Ross 308 strain unsexed chicks, seven-days old. Kernels, manually obtained from *J. curcas* seed, were defatted and boiled in order to obtain the treated jatropha kernel meal. This latter was used to replace the groundnut meal contained in a premix which was then incorporated in a commercial diet to warrant iso-nitrogenous and iso-caloric characteristics of the diets. The results revealed that animals that received the diet incorporating jatropha kernel meal had numerically higher live weight (272.6 vs. 259.3 g/animal) (P>0.05) and average daily weight gain (15.9 vs. 13.5 g/day/animal) (P>0.05) than the control ones, at the end of experiment. The average daily feed intake was a bit similar for the two groups (34.1 vs 34.5 g/day/animal) (P>0.05) as well as the feed conversion ratio (2.2 vs. 2.7 respectively for the jatropha group and the control group). The survival rate, at the end of the experiment, was 100% for the two groups of animals.

Keywords: broiler chicks; Jatropha curcas; defatted kernel; Senegal

Résumé

Effets de la substitution du tourteau d'arachide par l'amande de la graine de *Jatropha curcas* déshuilée et bouillie dans l'alimentation de poussins de chair au Sénégal

Cette étude a été conduite pour déterminer, dans un aliment croissance, l'effet de la substitution du tourteau d'arachide par du tourteau d'amande de *Jatropha curcas* ayant subi un traitement physico-chimique sur la viabilité, l'ingestion et les performances de croissance de poussins chair.

L'expérience de nutrition a duré sept jours avec vingt poussins non sexés de souche Ross 308, âgés de sept jours. Les amandes, obtenues manuellement à partir de graines de *J. curcas*, ont été déshuilées puis bouillies afin d'obtenir un tourteau d'amande de jatropha traité. Ce dernier a été utilisé pour remplacer le tourteau d'arachide contenu dans un prémélange qui a ensuite été incorporé à un aliment de type commercial pour maintenir les caractéristiques iso-azotées et iso-caloriques des régimes. Les résultats ont révélé que les animaux ayant reçu le régime contenant du tourteau d'amande de jatropha ont obtenu un poids vif (272,6 vs 259,3 g/ animal) (P>0,05) et un gain pondéral quotidien moyen plus élevés (15,9 vs 13,5 g/jour/animal) (P>0,05) que les animaux du groupe témoin. L'ingestion quotidienne moyenne était presque similaire pour les deux groupes (34,1 vs 34,5 g/jour/animal) (P>0,05) ainsi que l'indice de conversion alimentaire (2,2 vs 2,7 respectivement pour le groupe jatropha et le groupe témoin). Le taux de survie, à la fin de l'expérience, était de 100% dans les lots expérimentaux.

Mots clés: poussins de chair ; Jatropha curcas ; amande déshuilée ; Sénégal

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1. Introduction

Broiler chicken production remains one of the veritable ways of achieving sustainable and rapid production of high quality protein to meet the demand of the population (Apata & Ojo, 2000) especially in several developing countries like Senegal. In this country, the poultry sector has experienced significant development in its semi-industrial and industrial sector as well as in the family sector (Traoré, 2014). However, the dependence and the import of soybean have increased substantially the costs of poultry feed ingredients (Willems *et al.*, 2013). To overcome this, several non-conventional food sources have been proposed to be incorporated into poultry feed (Akinmutimi & Okwu, 2006; Dahouda *et al.*, 2009; Ayssiwede *et al.*, 2010; Diaw *et al.*, 2012; Kenis *et al.*, 2014). Based on that, *Jatropha curcas* was proposed to be incorporated into rations for poultry (Ojo *et al.*, 2013; Ojediran *et al.*, 2014; Nesseim *et al.*, 2015; Nesseim *et al.*, 2019).

This plant belongs to the *Euphorbiaceae* family, distributed all over the tropics and subtropics, able to grow on degraded soils (Heller, 1996). If most genotypes are toxic (Martinez-Herrera *et al.*, 2010), there are non-toxic genotypes which seeds can be consumed by humans (Makkar *et al.*, 1998b) and are also an excellent fish feed (Makkar *et al.*, 1999). The meal obtained after oil extraction is an excellent source of

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nutrients (Devappa & Swamylingappa, 2008) despite the presence of anti-nutrients and toxic components (Makkar et al., 2008). The toxicity of the seed from J. curcas was suggested to be due to phorbol esters, which are most concentrated in the kernel of the seed (He et al., 2011), have been shown to be the most important toxic molecules (Makkar et al., 1997; Becker & Makkar, 1998; Makkar et al., 1998a; Roach et al., 2012). Outside phorbol esters, jatropha meal contain, not only curcin (Asseleih et al., 1989) that is capable of inhibiting protein synthesis (Lin et al., 2003), but also anti-nutrients including trypsin inhibitor, phytate and saponins (Francis et al., 2002). The toxic genotype could be utilized as a fertilizer (Nithiyanantham et al., 2012) but efforts are under way to detoxify jatropha seed by removing phorbol esters or develop varieties that are deprived of this molecule so that the meal could be used as an ingredient in livestock diet (Makkar & Becker, 2009). Chemical de-oiling of jatropha kernel, followed by a physico-chemical treatment did not cause a complete removal of phorbol esters (Kumar et al., 2010b) and so many processing methods have been explored to detoxify meal of J. curcas with different levels of success. These include physical (Aregheore et al., 1998) and chemical (Haas & Mittelbach, 2000; Aregheore et al., 2003) methods; the combination of these two (Martinez-Herrera et al., 2006); and biological methods (Belewu & Sam, 2010; Joshi et al., 2011; Nesseim et al., 2019). Otherwise, curcin and trypsin inhibitor could interfere with physiological process of monogastrics causing severe growth depression (Palacios et al., 2004) but they may be removed by heat and biological treatment (Aderibigbe et al., 1997; Aregheore et al., 1998; Abou-Arab & Abou-Salem, 2010; Sumiati et al., 2012).

The objective of this study was to evaluate the impact of groundnut meal substituted by *J. curcas* kernel meal from Senegal that was subjected to combined chemical and thermal treatments in order to remove anti-nutritional compounds out of the product in a diet, on viability, ingestion and growth performance of broiler chicks.

2. Materials and methods

2.1. Location of the experiment

Experiment was conducted in *Ecole Nationale Supérieure d'Agriculture (ENSA)*, University of Thies (Senegal) just after the rainy season (November) with a temperature ranging from 26.9 to 37.8°C and a relative humidity ranging from 34 to 38%.

2.2. Collection and processing of jatropha seeds

Five hundred grams of mature and dry seeds of *Jatropha curcas* were collected from Dialacoto, geographical coordinates 13°19′0″ N and 13°18′0″ W, in the Tambacounda region (Senegal). The jatropha seeds cracked individually to remove the kernels. The kernels were washed with tap water to remove shells, drained properly and sun dried to a constant weight. These were milled into powder using WARING®-type speed blender with timer grinder and then defatted for 6 hours, by 20 g amounts each in a series Soxhlet-type extractor, using diethyl ether (boiling point, 60–80°C) as solvent.

The substrate, consisting of jatropha kernel meal, was spread

into a dish and boiled at 120°C for 30 minutes. The spent substrate, 239.6 g of boiled jatropha kernel meal (BJKM) was later used in the formulation of diet.

2.3. Diets preparation

Two broiler starter diets were formulated. The control diet (0BJKM) contained 2/3 of a complete starter commercial feed (SEDIMA S.A., Dakar, Senegal) for broiler chicks. This commercial feed was mainly composed of maize, cereal issues, soybean meal, peanut meal, fish meal, calcium carbonate, and vitamin-mineral complex. To formulate the control diet, 1/3 of a premix contained groundnut meal (160 g), corn (480 g), disodium phosphate (32 g), and calcium carbonate (32 g) was added to the commercial feed and the experimental diet (8BJKM) was formulated similarly but the groundnut meal of the previous mixture was replaced with boiled jatropha kernel meal (160 g) (Table I). These final mixtures were iso-nutrients for ME, CP, Ca, P, and Na, and in agreement with the recommendations of N.R.C. (1994).

Table I: Composition of diets incorporating J. curcas kernel meal

Raw materials (%)	0BJKM	8BJKM
Complete starter diet	68.0	68.0
Groundnut meal	8.0	-
BJKM	-	8.0
Maize	20.0	20.0
Phosphate disodium	2.0	2.0
Calcium carbonate	2.0	2.0
Total	100.0	100.0

Complete starter commercial feed (SEDIMA) composed of maize, cereals issues, soybean meal, peanut meal, fish meal, calcium carbonate, and vitaminmineral complex.

BJKM Boiled Jatropha Kernel Meal, *0BJKM* control diet, *8BJKM* diet incorporating 8% of boiled jatropha kernel meal.

2.4. Experimental animals and management

Twenty unsexed broiler chicks Ross 308 strain of seven-days old were used for the study. The birds were divided into two groups of ten chicks (control group -CG- and jatropha group -JG-) assigned to any of the two diets in a completely randomized design. Animals were kept in a naturally well-ventilated broiler chicken barn divided into two areas separated by a fence of 0.75 m in height. Each area was 2.25 m².

During the test, all day and all night, animals were heated by electric light (100 watts) ensuring luminosity and thermal comfort. The average temperature inside the experimental room was 31.6° C while the relative humidity was 37.7%.

Feed was weighted early in the morning; provided once a day and water was available *ad libitum*. Feed Refusals were collected and weighed the day after the distribution.

The study was carried out for seven days.

2.5. Data collection

Data were collected on feed intake, weight gain, feed conversion ratio and mortality.

The daily feed intake was estimated as the difference between the feed supplied and the feed rejected over 24 hours' period.



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The Average Daily Feed Intake (ADFI) was then estimated.

Birds in each replicate were individually weighed at the beginning and at the end of the experiment. This weighing was done in the morning, before the distribution of diets. The Average Daily Weight Gain (ADWG) was estimated.

An electronic scale with a weighing capacity of 50 kg and a precision of 0.01 g was used for all weight data.

The Feed Conversion Ratio (FCR) was determined as the feed intake per unit weight gain.

Mortality was recorded in each replicate and expressed as a percentage of the total number of birds in the replicate at the beginning of the experiment.

2.6. Chemical analyses

Samples of jatropha kernel meal, boiled jatropha kernel meal, control diet and diet incorporating the boiled jatropha kernel meal were analyzed. Dry matter (DM) was determined by oven-drying at 70°C for 15 hours, 90°C for 5 hours, and 102°C for 5 hours consecutively. Diets were analyzed for crude protein (CP; Method 954.01; AOAC, 1990), ether extract (EE; Method 920.39; AOAC, 1990) with petroleum ether solvent, ash (Method 942.05; AOAC, 1990), and crude fiber (CF; Method 962.09; AOAC, 1990).

The following values were calculated from those measured (Sibbald, 1976):

Organic matter (OM) = 100 - Ash

Non-Nitrogen Extract (NNE) = OM - EE - CP - CF

 $ME = metabolic energy (kcal/kg DM) = 3951 + (54.4 \times EE) - (88.7 \times CF) - (40.8 \times Ash)$

2.7. Statistical analysis

The presence or not of boiled jatropha kernel meal in diets is used to evaluate effects on ingestion and growth performance. Data generated were analyzed by one-way analysis of variance using the General Linear Model (GLM) procedure of SAS 9.1 software package. Significance was determined at p<0.05 and where this was indicated, Duncan's option of the same software was used to separate the means.

3. Results

3.1. Chemical composition of feed

Table II shows the analytical composition of the experimental diets. Both 0BJKM and 8BJKM showed almost similar value, with regard to the DM, CP, EE, ash and CF. The true metabolic energy of each diet was 3552.2 and 3366.5 kcal/kg DM for respectively 0BJKM and 8BJKM. Analytical results showed that diets were almost iso-proteinic and iso-energetic even if the metabolic energy of the control diet was higher than the boiled jatropha kernel diet.

Table II: Proximate analytical composition of the diets used during the experimentation

	DM	Chemical composition (% in DM)					ME	
	(%)	ОМ	СР	EE	CF	Ash	NNE	(kcal/kg DM)
JKM	91.6	90.1	61.8	5.0	8.2	9.9	15.1	3091.7
BJKM	97.4	90.6	54.0	4.3	7.5	9.4	24.8	3135.8
0BJKM	91.8	91.0	21.5	4.5	3.1	9.0	61.8	3552.2
8BJKM	90.1	89.2	22.4	3.5	3.7	10.8	59.5	3366.5

3.2. Growth performance and feed intake

Based on the visual observation during feeding time, palatability or acceptability of feed was good and the behavior of chicks was normal. There was no mortality during the entire experimental period.

Figure 1 shows synthetic body weight changes over the experiment. During the seven days, it was found that the control group showed a linear weight growth, evolving from 164.1 ± 28.8 g on day 8 (d8) to 259.3 ± 64.4 g on day 14 (d14). For the same period, animals that received the boiled jatropha kernel meal diet had the same profile, from 165.9 ± 11 g to 272.6 ± 35.4 g.



Figure 1: Weight performances of broiler chicks receiving or not 8% boiled *J. curcas* kernel meal

Figure 2 shows the daily individual feed intake of broiler chicks during the experimental sequence. No significant differences were observed in the feed intake of the two groups of animals $(34.5\pm4.9 \text{ g/d}/\text{animal} \text{ in CG vs. } 34.1\pm4.4 \text{ g/d}/\text{animal} \text{ in JG})$. For both groups of animals, a similar feed intake was noted. ADWG per animal did not change significantly for both groups regardless of the rate of boiled jatropha kernel meal incorporation, from 13.5 g/d/animal for CG to 15.9 g/d/animal for JG. The FCR presented the same mean values, 2.7 and 2.2, respectively for CG and JG without significant difference (P>0.05).



Figure 2: Growth performances of broiler chicks receiving or not 8% boiled *J. curcas* kernel meal.

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4. Discussion

Mechanical extraction of oil, from Jatropha curcas seeds, by means of a screw press is the method used generally in developing countries because of the simplicity of the equipment required (Eckart & Henshaw, 2012). This method makes it possible to recover more than 80% of the oil (Tambunan et al., 2012). However, solvent extraction could be regarded as the most ideal extraction method since it could recover 95-98% mass fraction of the available oil in the seed (Gübitz et al., 1999). In our study, because of the nondigestibility of hulls' fibers for monogastrics (Jørgensen et al., 1996), the jatropha seeds were manually shelled before being processed. The kernel obtained was crushed and completely de-oiled by the Soxhlet method. By choosing this method of de-oiling, our aim was to obtain a significant reduction of toxic compounds (Gandhi et al., 1995), allowing animals to ingest the jatropha meal. Our results were in line with those of Gübitz et al. (1999). But later studies (Makkar et al., 1998a; Martinez-Herrera et al., 2006) have shown that de-oiling did not allow this significant reduction.

An additional boiled treatment of jatropha kernel meal was done at 120°C for 30 minutes. The aim was to inactivate toxic and anti-nutritional compounds. Martinez-Herrera *et al.* (2006), by heat treatment in an autoclave (121°C for 20 mn), significantly inactivated trypsin inhibitor activity that is anti-nutritional factor but essentially inhibiting lectin activity which is considered to be another toxic factor in *J. curcas* seeds. In the same way, Abo El-Fadel *et al.* (2011) decreased the concentration of trypsin inhibitor and lectin by about 75 and 83% respectively. These results were in agreement with Haas & Mittelbach (2000) and Makkar *et al.* (2008) who reported also that heat treatment has a positive effect on reducing trypsin inhibitor and lectin concentration in *J. curcas* meal.

The chemical composition of jatropha kernel meal and boiled Jatropha kernel meal showed that the kernel meal contained almost the same ether extract as the boiled kernel meal of 5.0 and 4.3% DM respectively. The crude protein of Jatropha kernel meal was higher (61.8% DM) than boiled jatropha kernel meal (54.0% DM). This is an indication that during the soaking process, some nitrogenous substances in the meal were solubilized. Similar result was reported by Emenike *et al.* (2016) when they subjected raw jackbean to boiling process. However, the metabolic energy showed higher value for the boiled Jatropha kernel meal (3091.7 kcal/kg DM). Fakunle *et al.* (2013) obtained similar results with regard to metabolic energy; however, they obtained a higher crude protein level for the boiled jatropha kernel meal than the kernel meal.

Despite the BJKM incorporation, the daily feed intake per broiler chick that received this diet did not vary significantly compared to those who received the control diet (34.1 vs 34.5 g/d/animal). At the end of the experiment, the daily weight gain showed better values for the diet incorporating the boiled jatropha kernel meal than the control diet (15.9 vs 13.5 g/d/ animal), a better final weight (272.6 vs 259.3 g) and a better feed conversion ratio (2.2 vs 2.7) for the tested diet compared

to the control diet. These results are better than those obtained by Chandrasekar *et al.* (2009), Ojediran *et al.* (2014) who determined the growth performance of broiler chicks that fed from differently processed jatropha kernel meal. Antyev *et al.* (2017) obtained, even if they were lower than those of the control group, better final weight, daily weight gain and feed conversion ratio for broilers fed with diets incorporating either fermented jatropha kernel meal, or boiled jatropha kernel meal, compared to groups of animals that received diets incorporating kernel meal that had undergone other treatment.

Our results have, however, confirmed those of Alatise et al. (2014), Abozaid et al. (2016) and Musa et al. (2018) who evaluated on one hand, the effect of incorporating two levels of boiled jatropha kernel meal in place of soybean meal as a new source of protein on growth performance and carcass composition of Nile tilapia fingerlings, and an other hand, the growth response of Clarias gariepinus fingerlings using boiled jatropha kernel as a source of protein. Thus, Musa et al. (2018) by processing in boiling water, also have improved the nutritional profile of jatropha kernel meal and its utilization by African catfish Clarias gariepinus fingerlings in consonance with the studies of Adebayo et al. (2004) and Solomon et al. (2017). Our study, like that of these authors, has shown that a thermal regime for processing jatropha kernel meal ensured the improvement of the nutritional content and caused a significant reduction in most anti-nutritional factors of the unconventional feedstuff. These results do not correspond to those obtained by Jimoh et al. (2016) where the growth of clarias gariepinus reduced with the increase inclusion of cooked *jatropha curcas* due to the presence of anti-nutritional factors and poor digestibility of protein.

Our study did not allow measuring the levels of phorbol esters as well as curcin, but given the reaction of the chicks, without applying a chemical or biological detoxification treatment. It is possible that the presence of toxic factors was not sufficient to cause poor consumption and therefore poor weight gain.

The thermal regime for processing of *J. curcas* kernel meal ensured the improvement of the nutritional content and caused a significant reduction in most anti-nutritional factors of the unconventional feedstuff. In addition, the growth performance of chicks was improved, which was not the case in our previous study where we observed a growth depression and a poor feed conversion ratio in broiler chicks fed with jatropha kernel meal dry heat treated and incorporated in a diet (Nesseim *et al.*, 2017).

This study reveals the possibility of utilizing boiled jatropha kernel meal in the diet of chicks. Moreover, the treatment has even positively affected feed intake and weight and it had no impact on the viability of animals.

5. Conclusion

This study was the first field experiment on evaluation of boiled jatropha kernel meal in broiler chicks feeding in Senegal. The results showed that, after a total dehulling, a chemical deoiling with diethyl ether as well as a heat treatment, jatropha kernel meal did not affect viability, feed intake and growth of



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chicks. Further studies must be performed in order to confirm the use of the thermal processes to allow detoxification of *Jatropha curcas* meal.

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