Effects of Varying Fermentation Period on the Chemical Properties of Tropical Sikcle Pod (Senna obtusifolia) Seed Meal

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Abstract: A study was conducted to evaluate the proximate composition, amino acid profile and levels of antinutritional factors of *Senna obtusifolia* seeds subjected to varying fermentation periods. The seeds of *Senna obtusifolia* were naturally fermented for 0, 3, 5, 7 and 9 days, respectively. Each representative sample was analysed in triplicates for dry matter, crude protein, crude fibre, ether extract, nitrogen free extract, amino acids and levels of anti-nutritional factors using standard laboratory procedure. The results indicated an increasing trend as the fermentation period progresses for the protein (26.95 to 28.29%), ash (4.50 to 5.31%) and some amino acid content. Lysine and methionine for instance increased from 1.19 to 2.97% and 2.20 to 2.88%. As the fermentation period progresses, the crude fibre, nitrogen free extract, ether extract and anti-nutritional factors were observed to decrease. Crude fibre decreased from 11.17 to 4.23%, NFE 41.85 to 37.31% and ether extract 3.65 to 1.98%. Similarly, Tannins and oxalates for instance, decreased from 5.42 to 1.17 g/100g and 1.95 to 0.36 g/100g, respectively. It can be concluded that fermenting *Senna obtusifolia* seeds for up to 9 days has beneficial influence in improving the nutritional quality of *Senna obtusifolia* seeds. There is need to conduct further studies to investigate the effect of fermenting *Senna obtusifolia* seeds beyond nine (9) days.

[Augustine, C. Effects of Varying Fermentation Period on the Chemical Properties of Tropical Sikcle Pod (*Senna obtusifolia*) Seed Meal. *Academ Arena* 2018;10(1):49-54]. ISSN 1553-992X (print); ISSN 2158-771X (online). <u>http://www.sciencepub.net/academia</u>. 7. doi:<u>10.7537/marsaaj100118.07</u>.

Keywords: Sickle pod, chemical properties, fermentation period

Introduction

The utilization of lesser-known legumes will go a long way in addressing the feed crises that has engulfed the Nigerian livestock industry. Adegbenro et al. (2011) further buttressed the need to exploit some under-utilized seeds which could argument the costly conventional feed resources. One of such underexploited legumes is Senna obtusifolia seeds. Senna obtusifolia is a pantropical weed that belongs to the family leguminosae caesapinioideae. It is an erect bushy annual shrub that grows up to 90 cm tall and propagates through seed. The leaves are obovate and the flowers are yellow in colour (Akobundun and Agyakwa, 1998). Most seeds of legumes contain toxic components which could limit their utilization (Parul, 2014; Adebowale and Maliki, 2011). The chemical composition of the seeds as revealed by Ingweye et al. (2010) and Ismaila et al. (2011) indicated that they have good nutritional properties (29.54 and 18.46% crude protein) but also contain anti-nutritional factors (tannins 388.50 mg/100g, phytates 240.50 mg/100g and oxalate 83.25 mg/100g) which may adversely affect the performance of animals that consumed the seed meal. Processing treatments have been known to reduce antinutritional factors and improve their utilization

(Effiong et al., 2011; Tuleun et al., 2011; Emiola, 2013). Fermentation has been reported to modify some physical characteristics of cereals and legumes, increase the level of some nutrients, digestibility and bioavailability (Brown et al., 1998), decrease levels of anti-nutrients, increase nutrient density (Tomkins et al., 1987; Nnam, 1998), and imparts some antimicrobial properties (Mensah et al., 1990; Mensah et al., 1991). Fermentation holds promise as a food processing method that can be used to diversify the food uses of some exploited plant foods (Anthony and Babatunde 2014). Fermentation period has been documented to influence the chemical properties of seeds (Yashim et al., 2009 Adebowale and Maliki, 2011). At the moment, base-line information on the effects of varying fermentation periods on the proximate composition, amino acid profile and levels of anti-nutritional factors of Senna obtusifolia seeds seems to be meager. Therefore, more studies are needed to bridge such information gap. In view of the above, this study was designed to investigate the effects of varying fermentation periods on the chemical properties of Senna obtusifolia seeds.

Materials and methods

Seed collection and processing

The seeds were harvested from bushes around Mubi area of Adamawa State. The plant and seeds were authenticated at the Department of Biological Sciences, Adamawa State University, Mubi, Nigeria. The seeds were boiled, washed, drained and placed in an air tight container and allowed to naturally ferment for 3, 5, 7, and 9 days, respectively. At the end of each representative fermentation period, each representative sample was removed from the fermenting container and properly sundried, ground to meal and used for the chemical analysis.

Chemical analysis

The proximate composition of the seed meal and levels of the anti-nutritional factors were determined using the standard procedure of Official Association of Analytical Chemists [AOAC] (2004). The dry matter content was determined using the oven dry method and crude protein was determined using the Kjeldahl procedure. Soxhlet extraction method was used for the determination of ether extract, while the fibre content was evaluated using the trichloroacetic method. The ashing procedure was used to determine the ash content, while nitrogen free extract (NFE) was computed using the formula:

NFE = 100 - (% Moisture + CP + CF + EE + ASH)

Where:

CP = Crude Protein

CF = Crude Fibre

EE = Ether Extract

Similarly, the energy values were computed using the formula of Pauzenga (1985) which is expressed as $ME=37 \times \% CP + 81 \times \% EE + 35.5 \times \% NFE$

The amino acid profile was analyzed using High Performance Liquid Chromatograph (HPLC) Buck Scientific BLC 10/11 model equipment.

Data analysis

Data obtained were subjected to Analysis of Variance (ANOVA) and Duncan Multiple Range Test (DMRT) was used to separate the treatment means where significant difference occurred. The results were expressed as means with their standard error of the means of triplicate determinants.

Results and discussion

The proximate composition of Senna obtusifolia seeds subjected to varying fermentation periods is presented in Table 1. The proximate composition were significantly (P<0.05) affected by the different treatments. The crude protein was observed to increase with progressive increase in fermentation period. Fermentation period on the 9th day recorded the highest crude protein content. The crude protein content was observed to increase from 26.95 to 28.79%. An increase in the protein content after fermentation was due to the activities and increase number of lactic acid bacteria present during fermentation. The increase in protein contents agreed with the work of Afoakwa et al. (2004). This finding is also in line with the report of Adebowale and Maliki (2011) who obtained similar results for pigeon pea (Cajanus cajan) seed flour. They attributed such increase in protein content to net synthesis of protein by fermenting seeds which might have resulted in the production of some amino acids during protein synthesis (Marero et al. 1989; Uwagbute et al., 2000). Some workers (El Hag et al., 2002; Ali et al., 2003) reported that fermentation can be effectively used to improve the nutritional quality of cereal grains by increasing protein content and digestibility which is in line with the findings of this study.

The ash content was observed to significantly (P<0.05) increase with increase in the fermentation period. The ash content increase from 4.50 to 5.31% from 0 to 9 days. This result concurred with the findings of Adebowale and Maliki (2011) and Anthony and Babatunde (2014). The authors explained that the increase in the ash content after fermentation could be due to the incomplete utilization of minerals by the fermenting organisms during their metabolism.

The ether extract and nitrogen free extract (NFE) significantly (P<0.05) decreased as the fermentation period advances. The decrease in the ether extract was due to the breakdown of fatty acids and glycerol by lipolitic organisms during fermentation (Anthony and Babatunde, 2014). The NFE decreased from 41.85 to 37.31% with fermentation at the 9th day, recording the lowest value of 40.18%. This was due to the utilization of sugars in the seeds by the fermenting microbial mass which is in line with the report of Anthony and Babatunde (2014).

Fermentation period (days)						
Nutrients (%)	(T1)	T2(3)	T3(5)	T4(7)	T5(9)	SEM
Dry matter	90.72	90.33	90.27	90.18	90.23	21.43 ^{NS}
Crude protein	26.95°	26.49 ^c	27.34 ^b	27.19 ^{ab}	28.79^{a}	3.74*
Crude fibre	11.17^{a}	8.64 ^b	8.51 ^b	6.41 ^c	4.23 ^d	1.66*
Ether extract	3.65 ^a	3.42 ^b	3.39 ^b	2.52°	1.98 ^d	0.32*
Ash	4.50 ^b	4.54 ^b	5.01 ^a	5.11 ^a	5.31ª	0.44*
NFE	41.85 ^a	40.99 ^a	40.92 ^a	39.89 ^b	37.31 ^c	5.94^{*}
*Energy (MJ/Kg)	10.40	10.20	10.32	10.30	10.01	3.15 ^{NS}

	Table 1: Proxi	mate Comj	position of <i>Senna</i>	ı obtusifolia See	ds Subjected to	Varying Fermentation	1 Period
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a, b, c, d, e = Means in the same row with different superscripts are significantly different (P<0.05) * = Significant at 5% level of probability, SEM = standard error of mean

The crude fibre similarly indicated a decreasing trend with progressive increase in the fermentation period. Fermentation for 9 days recorded the least crude fibre content of 4.23% which is in agreement with the findings of Anthony and Babatunde (2014) who attributed such decrease to the enzymatic breakdown of fibre by the fermenting microbes which utilized them as carbon source and convert them to microbial biomass, thereby reducing the fibre content (Rainbault, 2001).

The amino acid profile (Table 2) also indicated a significant (P<0.05) increasing trend as the

fermentation period progresses. Fermentation was reported to cause biological enrichment of food substrate with protein, essential amino acids and vitamins (Skeinkraus, 1998) which is in agreement with the finding of this study. For instance, the highest lysine (2.97%) and methionine (2.88%) were recorded in the seeds fermented for 9 days. Net synthesis of protein by fermenting seeds resulted in the production of some amino acids during the protein synthesis (Marero *et al.* 1989; Uwagbute *et al.*, 2000).

 Table 2: Amino acid Profile (g/100g) of Senna Obtusifolia Seeds Subjected to Varying Fermentation Periods

Fermentation period (days)							
Amino acids	T1(0)	T2(3)	T3(5)	T4(7)	T5(9)	SEM	
Lysine	1.19 ^c	1.80 ^c	1.94°	2.37 ^{ab}	2.97 ^a	0.31*	
Methionine	2.20 ^c	2.31°	2.37°	2.48 ^{ab}	2.88 ^a	0.44*	
Threonine	2.25°	2.34 ^c	2.45 ^b	2.69 ^{ab}	2.78^{a}	0.10*	
Isoleusine	2.32 ^b	2.23 ^b	2.44 ^b	2.43 ^b	2.96 ^a	0.61*	
Leucine	3.59°	3.77 ^b	3.78 ^b	3.92 ^a	4.01 ^a	0.12*	
Alanine	0.97 ^e	1.15°	1.22 ^b	1.38 ^a	1.39 ^a	0.63*	
Phenylalanine	1.82 ^d	2.22 ^c	2.52 ^b	2.52 ^b	2.68 ^a	0.32*	
Valine	1.63°	2.34 ^b	2.39 ^b	2.45 ^a	2.65 ^a	0.71*	
Arginine	1.19 ^d	1.17 ^d	1.49°	2.16 ^b	2.59 ^a	0.13*	
Glutamic acid	0.94 ^e	1.35 ^d	1.43°	1.61 ^b	1.73 ^a	0.32*	
Proline	2.15 ^c	2.44 ^b	2.53 ^b	2.59 ^{ab}	2.69 ^a	0.46*	
Glycine	1.32 ^c	1.44 ^c	1.51 ^{bc}	1.60 ^a	1.67 ^a	0.92*	

a, b, c, d, e = Means in the same row with different superscripts are significantly different (P<0.05) * = Significant at 5% level of probability, SEM = standard error of mean

The levels of anti-nutritional factors (Table 3) showed a decreasing pattern as the fermentation period increases. Tannins and oxalates for instance, decreased from 5.42 to 2.02% and 1.95 to 0.28%, respectively. Anthony and Babatunde (2014) made similar observation for soya bean (Glycine max) flour; Magdi

(2011) for pearl millet; Lasekan and Shabnam (2013) for Rambutan (*Nephelium lappaceum*). This reduction may be due to some enzymatic reaction in addition, microorganisms' breakdown the carbon and nitrogen sources and use them for production of energy and their activity during fermentation (Hemingway, 1988). The

decrease in tannin has been associated to microbial activity during fermentation (Dhankher and Chauhan, 1987; Ikemefuna *et al.*, (1991; Elhag *et al.*, 2002) or to abstraction of hydride ions and rearrangement of the phenolic structures due to the acidic environment caused by the fermenting microbes (Towo *et al.*, 2006). The decrease in oxalate content was linked to reasons advanced by Simpson *et al.*, 2009) that a reduced PH caused by microbes during fermentation, can change insoluble oxalate bound ions to soluble oxalate content which will be used as energy source by oxalotrophic bacteria. The phytates level was similarly observed to decrease as the fermentation period progresses with fermentation period at the ninth day indicating the lowest level of phytate. This reduction might be due to low PH caused by fermenting microbes which has enhanced phytase activity resulting to lowering of the phytate content. The reduction in PH of fermented foods caused by the production of various organic acids such as lactic acid and acetic acid favours the activity of the enzymes phytates which is able to dephosphorylate phytate effectively. (Marfo *et al.*, 1990; Sanberg and Andlid 2002; Reale *et al.*, 2007; Abdeland *et al.*, 2011). This might be responsible for the reduction of the phytates content of *Senna obtusifolia* seed meal.

 Table 3: Levels of Ant-nutritional Factors (g/100g) of Senna obtusifolia Seeds Subjected to Varying Fermentation Periods

Fermentation period (days)						
Anti-nutrients	T1(0)	T2(3)	T3(5)	T4(7)	T5(9)	SEM
Oxalates	1.95 ^a	1.28 ^b	1.01 ^b	0.36 ^c	0.28 ^c	0.06*
Tannins	5.42 ^a	3.21 ^b	2.73 ^b	2.02 ^c	1.17 ^d	0.07*
Flavonoids	3.86 ^a	3.13 ^b	1.28 ^c	0.30 ^d	0.01 ^e	0.05*
Phytates	4.61 ^a	3.41 ^b	2.52 ^{bc}	1.71 ^d	0.25 ^e	0.01*
Saponins	2.37 ^a	1.78 ^b	1.90 ^b	1.94 ^b	0.35°	0.10*

a, b, c, d, e = Means in the same row with different superscripts are significantly different (P<0.05) * = Significant at 5% level of probability, SEM = standard error of mean.

Conclusion

It can be concluded that the proximate composition (protein and ash) and amino acid profile of Senna obtusifolia seed significantly increased with the in fermentation period. progressive increase Fermentation for up to nine days recorded the highest increase for protein, ash and amino acid content. A decreasing trend as the fermentation period progresses was observed for the nitrogen free extract, ether extract and levels of anti-nutritional factors. Based on the findings of this study, it is recommended that Senna obtusifolia seed can be fermented for up to 9 days without depreciation in nutritional quality. There is need to conduct further studies to investigate the effect of fermenting Senna obtusifolia seeds beyond nine (9) days.

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