ISSN 1119-7455

EFFECT OF MALTING PERIODS ON THE NUTRIENT COMPOSITION, ANTINUTRIENT CONTENT AND PASTING PROPERTIES OF MUNGBEAN FLOUR

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ABSTRACT

Mungbean grain has great potentials for product development, being rich in protein and other nutrients. This study evaluated the effects of malting periods on the nutrient composition, antinutrient content and pasting properties of malted mungbean flour. Flour samples were produced form mungbean grain malted for 24, 48, 72 and 96 h and assayed for proximate composition, selected mineral contents, vitamins A and B1, antinutrient contents, functional and pasting properties. Results showed that increasing the malting periods of mungbean grain significantly (p<0.05) increased the protein, ash, fibre and mineral contents but decreased fat and carbohydrate contents of its flour. Oil absorption capacities increased while bulk density, swelling and water absorption capacity decreased significantly (p<0.05) with increasing time of malting. Extending time of malting gave flours with reduced antinutrient, peak, trough, final and setback viscosities and pasting time but higher pasting temperature compared to unmalted flour. The study has shown that malting period modifies the nutrient composition, antinutrient contents and functionality of mungbean flour differently and invariably may affect the performance of the flour in product development. Mungbean grain malted at 72 h gave flour with highest quality establishing its optimal malting period at 72 h.

Key words: malting, nutrient composition, 'orarudi', pasting-properties, mungbean

INTRODUCTION

Legumes occupy an important position in the diets of people in developing countries because they are rich sources of protein, minerals and vitamins and can be used to supplement their staples which are majorly carbohydrate based. El Maki et al. (2007) reported that legumes are major contributors of protein in most African diets, but some of such legumes are unconventional and under-exploited. Such unconventional legumes have been associated with some bioactive compounds that can impact human health hence the need to study them. Among such unconventional legumes, is mungbean spp. locally known as 'orarudi'. Mungbean has high nutritional potentials (Mensah and Olukoya, 2007). Like most legumes, mungbean is rich in protein and essential amino acids, with exception of sulphur-containing amino acids (Khalil, 2006). Mungbean protein is rich in lysine but deficient in methionine (Anderson, 2007). There is a paucity of information on the use of mungbean malt in product formulation or value-added products in Nigeria particularly in infant foods. Mungbean grains like other legumes contain certain antinutritional factors (including polyphenols and

phytic acid) beside high viscosity which may limit their flour utilization in product formulation (Sandhul and Lim, 2007; Makumba *et al.*, 2016). It is therefore, pertinent to develop strategies to process the underutilized indigenous crops into flours of high nutritional quality and functionality that can be used in product development.

One of the simple traditional technologies adopted for improving the nutrient composition and functionality of plant foods is malting. Malting is controlled germination followed by controlled drying of the germinated kernels. Malting plays a significant role in promoting the development of hydrolytic enzymes, which are not present in nongerminated grains. It improves both the nutritional and functional properties of legumes (Mensah and Tomkins, 2003) and allows preparation of low-bulk foods through elaboration of amylases resulting in reduced viscosity of the gelled germinated starch (Brandtzaeg et al., 1981; Kulkarni et al., 1991). Low bioavailability of nutrients arising from the presence of antinutrients such as phytate, polyphenols, and oxalate, could limit the quality of predominantly plant-based diets. Uppal and Brains

(2012) reported that malting could significantly raise the bioavailability of nutrients. Malleshi and Klopfenstein (1996) also reported that malting of grains increased protein and carbohydrate digestibility, enhance some of their vitamin contents, reduce antinutritional factors and improves their overall nutritional quality. The nutritional composition and antinutritional factors of mungbean seeds (*Phaseolus aureus*) as affected by some home traditional processes was also reported by Mubarak (2005).

Presently, utilization of malted legume flour like its cereals counterpart is gaining height in product development in developing countries and establishing their optimal malting period to obtain best flour becomes necessary. Kaur et al. (2018) reported that starches from different cereal exhibit different pasting and functional properties, however flours developed from legumes malted at different periods may perform differently in product development also and needed some investigation. According to Onwurafor et al. (2017, substitution of cereal flour with malted legume flour gave products of higher quality and better functional properties. Savelkoul et al. (1992) reported that the effect of germination depends on the conditions and duration of germination process. Research on the malting of legumes such as bambara groundnut, soyabean among others have been done. However, there is dearth of information in this regard of this specie of mungbean grain 'orarudi', necessitating research effort. For efficient utilization of the flour of malted mungbean flour, there is need to study the nutrient composition, antinutrient content as well as pasting properties in order to know the food system it can be applied in. Hence, this study evaluated the effects of malting periods on the nutrient composition, antinutrient content and pasting properties of mungbean flour.

MATERIALS AND METHODS

Mungbean grains (*Vigna radiata*) were purchased from Orba market in Nsukka Local Government Area, Enugu State, Nigeria. Identification of the crop was done in the Department of Crop Science, University of Nigeria, Nsukka.

Processing of Mungbean Grain into Flour

Five lots of sorted mungbean grain (200 g each) were weighed into porous bags (25 cm \times 45 cm) and steeped in water for 3 h, air rested for 90 min. and re-steeped in fresh tap water for 3 h (a modification of the two-step wet-steep method of Etok-Akpan and Palmer (1990). The steeping schedule was based on the time for maximum water absorption characteristics of the mungbean seed (Umunnakwe, 2012). One lot of mungbean seeds was wet dehulled. The second, third, fourth

and fifth lots of 200 g of grain each was spread in a dark room to germinate for 24, 48, 72 and 96 h, respectively, during which they were, turned every 24 h (30 ± 0.2 °C). The samples were moistened on alternate days by dipping the malting bags containing the germinating grains in water for 30 sec. One lot (bag) was picked at interval of 24, 48, 72 and 96 h. The wet dehulled sample and green malts were dried at 50 °C for 12 h and then separated from the sprouts and hulls by abrasion between the palms followed by winnowing. Subsequently, each lot was milled using a Bentall attrition mill (Model 200 L090, E. H. Bentall U.K), sieved through 200 μ m sieve and stored in polyethylene bags at 4 °C until use for analyses.

Determination of Proximate Composition, Minerals, Vitamins and Antinutrients

Mungbean malt flour was analyzed for proximate composition (moisture, crude protein, fat, fibre and ash) using AACC (2000) methods. Protein content of the samples was calculated using the formula, protein=nitrogen \times 6.25. Carbohydrate was estimated by difference (100 - % protein +% ash + % fat + % crude fibre + % moisture) and the values converted to g/100g sample. Each analysis was carried out in three replications. Calcium, iron, zinc, phosphorous, potassium, sodium, magnesium and copper contents of the malt flour were determined using inductively coupled plasma atomic absorption spectrophotometer (AACC, 2000). The processed flours were analyzed for provitamin A content according to the methods described by Arroyave et al. (1982) and converted to retinol equivalent. Vitamin B1 was determined according to the methods of AOAC (2010). Tannin, oxalate and phytate contents of the processed flours were determined according to the methods described by Burns (1971), Fassett (1973) and Latta and Eskin (1980), respectively.

Determination of pH, Titratable Acidity, Functional and Pasting Properties

The pH and titratable acidity were carried out using the methods described by AACC (2000). Swelling capacity was determined according to the methods described by Takashi and Sieb (1988) while water absorption and oil absorption capacity of the flour was determined by methods of AACC (2000). The bulk density was determined by the methods described by Okaka and Potter (1979). Pasting properties; peak (PV), trough (TV), breakdown (BD), final V(FV) and setback (SB) viscosities, peak time and pasting temperature of malted and unmalted flours were determined using Rapid Visco-Analyzer (Newport Scientific, PTY.LTD, Australia) connected to a computer (PC) with WindowsTM operating system via a USB port according to the methods described by Newport Scientific Method (1998). The moisture content of the flour was first determined to obtain the correct sample weight and amount of water required for the test. An aqueous suspension of sample was then made by mixing 3 g sample in 25 mL of distilled water and spun at 75 rpm. The temperature-time conditions included a heating step from 50 to 95 °C at 6 °C /min (after an equilibrium-time of 1 min. at 50 °C), a holding phase at 95 °C for 5 min., a cooling step from 95 to 50 °C for 2 min. Viscosities were expressed in rapid viscosity units (RVU).

Statistical Analysis

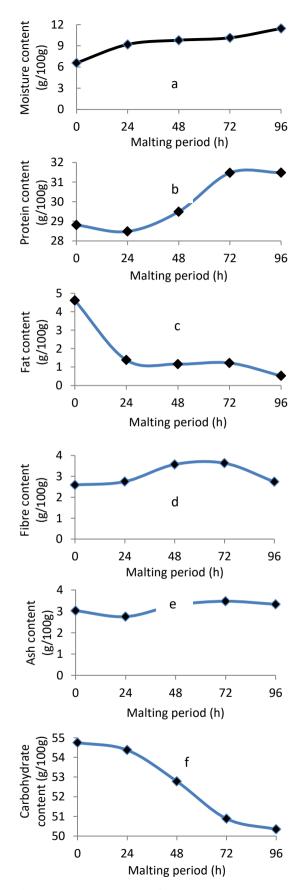
Data were subjected to one-way analysis of variance (ANOVA) (Steel and Torrie, 1980). Mean separation was done by Duncan New Multiple Range Test using SPSS version 16.00 software, with p set at < 0.05.

RESULTS AND DISCUSSION

Proximate Composition of Mungbean Malted at Different Periods

Figures 1a-f show the change in proximate composition of the flours of mungbean seeds malted at different periods. Moisture content increased as the malting period increased (Figure 1a). This observation is attributable to hydration of the seeds during steeping and germination. Also, since starch and proteins are broken down during germination, the sorption isotherm will be changing so that it may attract more water. Oluwole et al. (2012) attributed similar observation during malting of sorghum and maize grain to prolonged addition of water. A similar increase was observed in protein content of the flour with increase in malting period (Figure 1b). However, the protein content decreased after 72 h. The breakdown of complex proteins into simpler forms and/or reduction of other components such as carbohydrates and fats in the malts as reported by other researchers (Mubarak, 2005) may have given rise to the observed effect on protein content with increase in malting period. The protein value observed in the malts were higher than values from processed sunflower seeds (Adesina, 2018).

Progressive reduction of fat content of mungbean flour was observed as malting period was increased (Figure 1c). Flour sample malted for 96 h had the lowest fat content. The lower fat content of malted sample compared to the value of the unmalted flour (Figure 1c) agrees with the findings of Mubarak (2005). The use of fat as an energy source and increased activities of the lypolytic enzymes during malting which hydrolyzed fats to fatty acids and glycerol may explain the decrease in fat content. This shows that increase in malting period increases consumption of fat and the activities of lypolytic enzyme. However, an increase in fat content of soybean germinated for 1-6 days has been reported (Kayembe and Jansen van Rensburg, 2013).



Figures 1 a, b, c, d, e and f: Proximate composition (g/100g) of mungbean malted at different periods

Both fibre and ash contents of the mungbean malt (Figure 1 d) significantly ($p \le 0.05$) increased with malting period with highest values in samples malted for 72 h (Figure 1e). The result agrees with Chikwendu (2003) who attributed ash increase to endogenous hydrolysis of complex organic compounds which released more nutrients. On the contrary, the findings of Oluwole et al. (2012) and Muhammed et al. (2012) reported decreases in ash content during malting of sorghum and barley due to differences in seed composition. Lowest content of carbohydrate was observed in 96 h malt indicating that carbohydrate decreased with malting period. The decrease in carbohydrate and fat may have caused the increase in ash content of the samples as the malting period increased. Carbohydrate content decreased with malting period probably due to carbohydrate use as energy source for sprouting (Mubarak, 2005). Kirk-Othmer (2007) reported that malting affected carbohydrate molecules in germinating grains through the action of amylase whereby the carbohydrates were reduced to maltodextrins and low molecular weight sugars.

Minerals, Vitamin A and B1 Contents of Mungbean Malted at Different Periods

The values of calcium and iron increased as the malting period increased (Table 1). Malting for 72 h showed highest increase in calcium contents, the value finally decreased in the 96 h malted sample. The increase in mineral level with malting period increase may be due to concentration. Calcium and iron contents increase during malting of beans were earlier reported by Muhammad (1990). The K, P, and Mg contents of mungbean malt increased also as the malting period increased with 72 h malt having the highest values (1376.78, 312.99 and 187.31 mg/100g for K, P and Mg, respectively).

In contrast, zinc and sodium contents decreased for the first 48 h and then increased as malting period was increased to 72 h. The unmalted sample had higher zinc content (5.16 mg/100g) compared to 24, 48, and 96 h malt (4.71, 4.60 and 3.82 mg/100g, respectively) which increased at the 72 h of malting. The decrease in zinc content at the initial stage (24th and 48th h) could be due to the use of zinc in cell reproduction and tissue growth (Scherz and Kirchhoff, 2006). Zinc deficiency causes growth retardation and inadequate sexual development in human (Wardlaw and Kessel, 2002). The result indicates that mungbean malt is a good source of iron and zinc being significant at this time that iron and zinc deficiencies are among public health concern. The vitamin A as retinol equivalent (Re) and vitamin B₁ contents of the processed mungbean malt sample are also shown in Table 1. Vitamin A content of mungbean samples increased as the malting period increased. The non malted flour was found to have the least vitamin A content (43.7 µgRE/100g) while mungbean sample malted for 96 h had the highest vitamin A content (163.8 µgRE/100g). The increases in vitamin A (as retinol equivalent) with increasing malting period suggested that higher synthesis and/extractability of β -carotene occurs as the time of malting increased. Vitamin B₁ follows the same trend. Vitamin B_1 (thiamin) content of the processed mungbean flour ranged from 4.6 to 10.05 mg/100g. It was observed that vitamin B₁ increased as malting period increased. Non-malted flour had the least vitamin B_1 content while mungbean sample malted for 96 h had the highest vitamin B_1 content. Germination has been reported to increase vitamin B₁ (Nzelibe and Onyeniran, 2001).

Anti-nutrient Contents of Mungbean Malted at Different Periods

Length of malting affected the quantity of tannin, phytate and oxalate in the mungbean malt flour positively (Figure 2). The tannin, phytate and oxalate contents of mungbean malt flour decreased as the period of malting was increased. The decrease observed in these parameters was significant (p < 0.05). However, a marginal increase in tannin content was observed in the 72 and 96 h malted samples. The decrease in phytate contents could be attributed to increased activity of phytase due to increase in malting periods that progressively degraded phytic acid. These results corroborate Pawar and Machewad (2006) who attributed the reduction of phytate to phytic acid degradation by phytase synthesized during the process.

Table 1: Changes in selected mineral contents of mungbean during malting

		<u> </u>				
Parameter (mg/100g)	Malting period (h)					
	0	24	48	72	96	
Calcium	98.46 ^b ±0.69	$100.52^{b} \pm 0.01$	101.74 ^c ±0.02	122.00 ^e ±0.79	102.98 ^b ±0.91	
Zinc	5.16°±0.03	$4.71^{\circ} \pm 0.12$	$4.60^{b} \pm 0.00$	$5.31^{d} \pm 0.00$	$3.82^{a} \pm 0.00$	
Iron	$8.06^{b} \pm 0.04$	$7.88^{a} \pm 0.03$	$7.84^{a} \pm 0.06$	$11.02^{d} \pm 0.01$	$10.70^{\circ} \pm 0.01$	
copper	$0.73^{a} \pm 0.00$	$0.73^{a} \pm 0.00$	$0.78^{a} \pm 0.01$	$0.92^{\circ} \pm 0.02$	$0.79^{ab} \pm 0.00$	
Sodium	$5.47^{b}\pm0.17$	5.26 ^b ±0.03	5.11 ^a ±0.01	$7.07^{d} \pm 0.020$	5.63°±0.04	
Potassium	1147.80 ^a ±1.05	1148.89 ^a ±0.03	1270.21 ^b ±0.00	1376.78 ^d ±0.01	$1305.00^{\circ} \pm 0.01$	
Phosphorus	$261.06^{a} \pm 0.01$	262.56 ^a ±0.07	274.88 ^b ±0.05	312.99 ^c ±0.54	$288.87^{b} \pm 0.06$	
Magnesium	147.03 ^a ±0.507	147.54 ^a ±0.06	$166.42^{b} \pm 0.56$	187.31 ^c ±0.50	165.15 ^b ±0.00	
Vitamin A (µgRE/100g)	43.7 ^a ±0.02	$54.6^{a} \pm 0.00$	$65.5^{b} \pm 0.00$	$109.2^{d} \pm 0.00$	$163.8^{\circ} \pm 0.00$	
Vitamin B1(mg/100g)	$4.6^{b} \pm 0.01$	$5.3^{b} \pm 0.00$	$6.7^{\circ} \pm 0.00$	$7.7^{d} \pm 0.02$	$10.05^{e} \pm 0.01$	

Values are means $(n = 3) \pm SD$. Means on the same row carrying different superscripts are significantly (p < 0.05) different

Functional Properties of Mungbean Malt Flour Malted at Different Periods

There were significant (p < 0.05) differences in the water absorption capacity of all the samples (Table 2). Increasing the malting time significantly (p < 0.05) decreased the water absorption capacity and bulk density (BD) of the sample. Decrease in water absorption capacity as malting period increased may be due to hydrolysis of starch polymer (Adebowale et al., 2005) caused by malting effect or variation of protein content of mungbean malt that denatured during processing; this may affect the amount of water absorbed (Omeire et al., 2014). Protein may be hydrolyzed during germination to release free amino acids and these would bind water more effectively.

Swelling capacity of the mungbean flour samples varied from 0.553 to 0.763 g/ml and significantly (p < 0.05) decreased with increasing malting period. Differences in the friability of the materials caused by differences in malting period gave rise to variation in bulk density. However, there was marginal decrease in swelling capacity until the 48^{th} h, then significant (p < 0.05) decrease was seen in 72 and 96 h malted flour. Swelling capacity influenced other functional properties of starchy product which in turn determine their suitability in product development (Amajor et al., 2014). Generally, the low swelling capacity observed in the study could be due to malting effect caused by high amylose content of mungbean malt flour. Adebowale et al. (2005) noted that high amylose content could result in low swelling index. Increasing malting period may have effect on the starch component of the flour which may be caused by high amylose content of mungbean malt as noted earlier thus leading to reduction of swelling capacity. On the contrary, Kaur et al. (2018) observed higher swelling capacity in isolated cereal starches implying that cereal starch swells better than legume. Difference in structure of starch granules between cereals and legumes and /or pure starch in relation to whole grain flour may cause flour to swell differently. In contrary, oil absorption capacity increased with malting periods. Oil absorption capacity indicates the ability of the sample to retain flavor and improve mouth-feel (Kinsella, 1976). From the study, flour malted for longer period may have better flavor retention and mouth feel when developed into product.

Table 2: Some functional properties of mungbean malt flour

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Malting	g WAC	Bulk	Swelling	OAC
Period	(ml/100g)	density	capacity	(ml/100g)
(h)	-	(g/ml)	(ml/g)	-
0	153.00 ^d ±0.11	$0.763^{d} \pm 0.01$	0.763°±0.62	164 ^a ±0.03
24	147.00 ^{cd} ±0.06	0.732°±0.02	0.741°±0.87	188 ^a ±0.03
48	140.00 ^{bc} ±0.10	$0.706^{bc} \pm 0.01$	0.706°±0.13	173 ^a ±0.12
72	133.33 ^{ab} ±0.06	$0.693^{b}\pm0.01$	0.693 ^b ±0.07	233 ^b ±0.12
96	129.00 ^a ±0.05	0.627 ^a ±0.03	0.553 ^a ±0.02	247 ^b ±0.23
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Values are means $(n = 3) \pm SD$. Means on the same column carrying different superscripts are significantly ($p \le 0.05$) different; WAC - water absorption capacity; OAC - oil absorption capacity

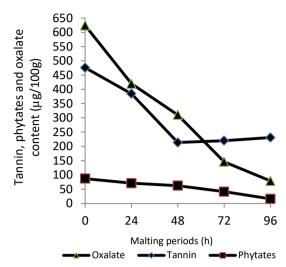
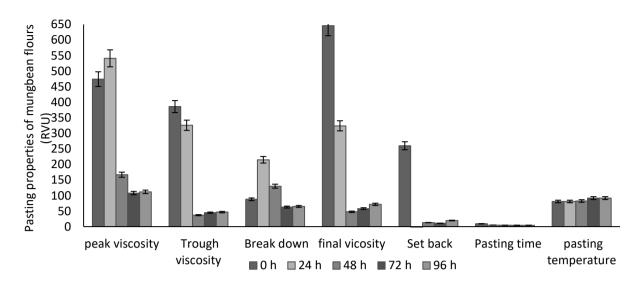


Figure 2: Anti-nutrient contents of mungbean malted at different periods

Pasting Properties of Mungbean Malt Flour Malted at Different Periods

Figure 3 shows the pasting properties of flour from mungbean grain malted at different periods. Unmalted mungbean malt flour was characterized with high peak viscosity (541 RVU), trough viscosity (386 RVU), final viscosity (647 RVU) and set viscosity (260 RVU) compared to malted samples. Extending malting periods significantly (p < 0.05) reduced the aforementioned properties. Bulk density viscosity was highest in 24 h malted sample. Increasing the malting periods resulted to significant (p < 0.05) decrease in peak viscosity (PV), trough viscosity (TV), final viscosity (FV) and setback (SB). The sample malted for 72 h had the lowest peak viscosity value.

The variation in peak viscosity values of flour samples obtained at different malting periods indicated that malting period confers different degree of gelatinization and affects the amylose content of the flour (Ayinde et al., 2012). Longer malting period may confer higher gel strength and tendency to the flour during cooking or reconstitution but also low reconstituted gel stability contrary to the report of Kaur et al. (2018) on starch isolated from six major cereal with higher peak viscosity (PV) values. The low final viscosity observed in the samples malted for 72 and 96 h indicated that extending period of malting may cause samples to form a low viscous paste rather than thick gel on cooking thus bring about thinning effects desired in certain products. The setback (SB) values suggest that increasing malting period would result to low retrogradation tendency and syneresis of the flour samples during freeze-thaw process. It was observed that the pasting temperature (PT) increased from 80.95 °C in non-malted flour sample to 91.90 °C. Increasing the malting period increased the pasting temperature and decreased the



pasting time. Inability of the flours to absorb water easily and swell due to malting periods could have increased the PT. On the contrary, Kaur et al. (2018) reported lower pasting temperature of most isolated pure cereals and attributed it to high water holding capacity of the starches. The PT was reported to be affected by starch concentration. Generally, high gelatinization temperature was observed in this study compared to values reported in literature for cereal starches and composite flour by some earlier researchers (deMan, 1999; Adeguwa et al., 2012; Onwurafor et al., 2017; Kaur et al., 2018). This indicated that the semi-crystalline nature of starch granules in mungbean flour malted at different periods will lose their birefringence differently and at higher temperature than those of pure cereal starches (Kaur et al., 2018). Unmalted sample had the highest cooking time, an indication that higher energy is needed to break the intermolecular bonds in starch granules of this flour to achieve gelatinization compared to malted sample (Kaur and Singh, 2006). This implies that increasing malting period may cause reduction in cooking time of flour, hence reduction in energy cost.

CONCLUSION

Malting of mungbean grains for different periods confer different properties to its flour in terms of nutrient composition, functional and pasting properties. Malting of mungbean grains up to 72 h increased the pasting temperature but reduced the cooking time. The optimal malting period for the mungbean grains was 72 h which had the highest content of most nutrients. The findings of this study could be applied at the rural community level to improve the nutrient and functional properties of the flours for use in product development.

ACKNOWLEDGEMENT

The authors wish to acknowledge the USDA through the Norman E. Borlaug Fellowship Programme in Pennsylvania State University, USA that gave access to some facilities for part of this research work.

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