

2 **Proximate composition, Functional and Sensory Properties of Pearl Millet, Soy flour**
3 **and Baobab Fruit Pulp Composite flour as a Complementary Food**

4
5 **Abstract**

6 **Aim:** to evaluate the proximate composition, functional and sensory properties of a
7 complementary food from pearl millet, soy flour and baobab fruit pulp composite flours.

8 **Study Design:** A complementary food was produced from Pearl millet, soy flour and baobab
9 fruit pulp powder (BFP) of various proportions (10, 20, 25 and 30%). Proximate (protein,
10 ash, moisture, fibre, fat, carbohydrate and energy value) composition, functional (Bulk
11 density, gelation capacity, swelling index, water absorption capacity and oil absorption
12 capacity) properties and sensory (appearance, flavour, texture and overall acceptability)
13 attributes were determined.

14 **Results:** The results of proximate composition showed that Moisture content ranged from
15 10.09 – 10.98, Protein content ranged from 9.80 – 24.25, Fat content ranged from 4.94 –
16 16.65, Carbohydrate content ranged from 43.11 – 71.03, Fibre content ranged from 3.37 –
17 15.67, Ash content ranged from 2.59 – 2.87% and Energy value ranged from 367.78 – 423.69
18 Kcal. The functional properties showed that Water Absorption Capacity ranged from 2.70 –
19 2.91, Oil Absorption Capacity ranged from 1.90 – 2.72, Bulk Density ranged from 0.69 –
20 0.71, Swelling Index ranged from 0.68 – 1.04g/ml and Gelation Capacity ranged from 5 –
21 10% of the complementary food samples. The sensory attribute also revealed that the
22 complementary food samples proved to be of good quality but the controlled sample (A) was
23 most preferred by the panellist.

24 **Conclusion:** The addition of baobab fruit pulp (BFP) to pearl millet and soybean flour, in
25 turn increases the fibre, ash and carbohydrate contents of the complementary foods. The
26 functional properties also improved with addition of baobab fruit pulp levels. This
27 improvement could be noticed in water absorption capacity, oil absorption capacity, bulk
28 density and swelling index. The sensory attributes indicates that the baobab fruit pulp
29 samples competes very well with the control (A) sample. However, sample A was most
30 preferred by the panellist.

31 **Keywords:** Baobab Fruit Pulp (BFP), Pearl Millet, Soybean, Complementary Food

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35 INTRODUCTION

36 Malnutrition is responsible, directly or indirectly, for over half of all childhood deaths.
37 Infants and young children are at increased risk of malnutrition from six (6) months of age
38 onwards, when breast milk alone is no longer sufficient to meet all nutritional requirements
39 and complementary feeding needs to be started. Complementary foods are often of lesser
40 nutritional quality than breast milk. In addition, they are often given in insufficient amounts
41 and, if given too early or too frequently, they displace breast milk. Complementary foods are
42 food other than breast milk or infant formula such as solid, liquid and semi-solid food
43 materials which are introduced to infants to provide nourishment (Anigo *et al.*, 2010). Gastric
44 capacity limits the amount of food that a young child can consume during each meal.
45 Repeated infections reduce appetite and increase the risk of inadequate intakes. Infants and
46 young children need a caring adult or other responsible person who not only selects and
47 offers appropriate foods but assists and encourages them to consume these foods in sufficient
48 quantity (WHO, 2001). It is common knowledge that breast milk is the best food for infants
49 during their first six (6) months of life. Breast milk contains all the essential nutrients and
50 immunological factors an infant requires to maintain optimal health and growth. It also tends
51 to protect infants against upper respiratory infection and diarrhea which are the chief causes
52 of infant and child morbidity and mortality (Cristina *et al.*, 2004 and Solomon, 2005).
53 However, at an early age of six (6) months and above, the weight of the child is expected to
54 double which breast milk alone at this point may not be sufficient for the child's nutritional
55 and growth needs. The adoption of recommended breast feeding and complementary feeding
56 practice and access to the appropriate quality and amount of foods are essential component of
57 optimal nutrition for infant and young children (Anigo *et al.*, 2010). Several factors tend to
58 contribute to the vulnerability of children (infants) during the complementary feeding period.
59 These factors may include; low nutritional quality of complementary foods which most times
60 are provided in insufficient amount to the child (WHO, 2002; Anigo *et al.*, 2010). In recent
61 years, many important advances in breast feeding promotion have been made but
62 unfortunately the same may not be said for complementary feeding (PAHO/WHO, 2003).
63 Some nutritional importance of the raw materials used The dried baobab fruit powder
64 contains about 12% water and various nutrients including carbohydrates, dietary fibre, B-
65 vitamins, calcium, magnesium, potassium and iron. The fruit is 100% natural and known for
66 its high content of vitamin C, pro-vitamin A, vitamin E, essential amino acids and calcium.
67 All of this anti oxidant are extremely important in human nutrition. Soybean also contains the
68 followings; Protein and oil makes up about 60% of the soybean and about one third consist of

69 carbohydrates, including polysaccharides, starchyose (3.8%), raffinose (1.1%) and sucrose
70 (5%), Phosphatides, sterols and other constituents. A variation ranging from 13.9 – 23.2% in
71 oil and 32.4 – 50.2% in protein has been recorded. The variation in protein and oil content in
72 soybean is due to the locality where the beans are grown. Literature reviewed that oil, sugars
73 and other non-protein components were affected mostly by changes in the protein content. An
74 increase in the protein content leads to a significant decrease in the non-protein constituents
75 such as oil, sugar and pentosans. Pearl millet contains 5.8 – 20.9% protein, 63.1 – 78.5%
76 carbohydrate, 1.4 – 2.6% soluble sugars, 1.1 – 1.8% fibre content and 4.1 – 6.4% fat content.
77 According to research in Georgia, pearl millet is 8 – 60% higher in protein and 40% higher in
78 lysine than is feed corn. Pearl millet is much lower in tannin than sorghum. Millet is high –
79 energy, nutritious food, especially recommended for children, convalescents and the elderly.
80 Several food preparations are made from millet which differs between countries and even
81 between different parts of a country. These consist primarily of porridge or pancakes-like flat
82 bread. However, because wholemeal quickly goes rancid, millet flour can be stored only for
83 short periods (F.A.O, 2007). Pearl millet is rich in B group vitamins, potassium, phosphorus,
84 magnesium, iron, zinc, copper and manganese. It is a gluten free grain and the only grain that
85 retain its alkaline properties after being cook which is ideal for people wheat allergies.
86 Commercial baby food formulae are made to the highest microbiological specification and
87 are formulated to meet the nutritional requirement of babies. They are designed to
88 complement normal family and more appropriate than adult convenience foods. Commercial
89 baby foods provide energy, protein, carbohydrate and fats. It also contain controlled amount
90 of fibre, sugar and salt. Vitamins and minerals such as vitamin C and Iron are essentially
91 added to the required amount. This research is therefore aimed at improving the quality of
92 complementary food through the supplementation of Baobab Fruit Pulp with other cereal e.g
93 pearl millet and Legumes such as soybean improve the nutritional quality of infant formula.
94 This research therefore aims to improve the quality of complementary food through the
95 supplementation of Baobab Fruit Pulp with other cereal e.g pearl millet and Legumes such as
96 soy flour to improve the nutritional quality of infant food.

97

98

99 MATERIALS AND METHODS

100 Materials

101 The food commodities used for this research were pearl millet (*Pennisetum glaucum*),
102 soybean (*Glycine max. L*) and Baobab fruit pulp (*Adansonia digitata*). Soybean and pearl
103 millet were purchased from North Bank market Makurdi, were brought to the University of
104 Agriculture Makurdi seed research centre for identification. Baobab fruit pulp powder was
105 obtained from Lafia Market in Nasarawa State. Nigeria

106

107 Pearl Millet Flour Preparation

108 The process of flour preparation as shown in fig 1 consists of dry cleaning of the pearl millet
109 i.e winnowing etc. The kernels were thereafter dehulled after mild wetting using rice
110 dehuller. The grains were then washed and dried in a convection hot air laboratory oven
111 (MODEL TT-9053 (Techmel and Techmel) at 50⁰ C for 24 hrs to 14% moisture content. The
112 dried grain was milled using a single disk attrition mill and sieved through a 455µm screen
113 laboratory sieve (MODEL STMN 2-CO402 JAPAN) and the under flow was used for the
114 research (Filli, *et al* 2012).

115 Soy Flour Preparation from

116 The method of Filli *et al*, (2012) was adopted as shown in fig 2. Soybean seeds were steeped
117 in clean tap water at 28⁰C for 24hrs in a plastic bowl. The kernel was therefore dehulled using
118 the traditional pestle and mortar. The grains were then washed and the hulls removed. After
119 which it was dried in a convectional laboratory hot air oven (MODEL TT-9053 (Techmel) at
120 50⁰C for 24hrs to 14% moisture content and the mass was winnowed to remove the
121 remaining lighter material using trail. The dehulled soybeans kernels were ground in a
122 laboratory disc attrition mill to fine flour. The flour was sieved through a 455µm screen
123 laboratory sieve (MODEL STMN 2-CO402 JAPAN) and the under flow was used for further
124 use.

125 Baobab Fruit pulp Flour Preparation

126 Baobab pods were cracked using a hammer. The pulp and seeds were transferred into a
127 ceramic mortar and it was pounded using a pestle until all the pulp was separated from the
128 seed. The pulp was sieved through a 455µm screen laboratory sieve MODEL STMN 2-

129 CO402 JAPAN to remove the fibrous materials from the pulp and the under flow was used
130 for further use as shown in fig 3

131

UNDER PEER REVIEW

132

133

Pearl millet

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135

Cleaning/washing

136



137

Oven drying (50⁰C for 24hrs)

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139

Weighing

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141

Toasting in microwaving (80⁰C for 15 min)

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143

Cooling

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145

Winnowing

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147

Milling

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149

Sieving (455μm)

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151

Flour

152



153

Packaged and store

154

Fig 1: Flow chart for the production of pearl millet flour.

155

Source: (Filli, 2012) with slight modification

156

157

Soybeans

158



159

Sorting

160



161

Cleaning

162



163

Blanching (60⁰C for 20 – 25 min)

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165

Dehulling by hand rubbing

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167

Removal of hulls by floatation

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169

Oven drying (55⁰ C for 24hrs)

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171

Toasting in microwaved (75⁰ C)

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173

Milling

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175

Sieving (455µm)

176



177

Flour

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179

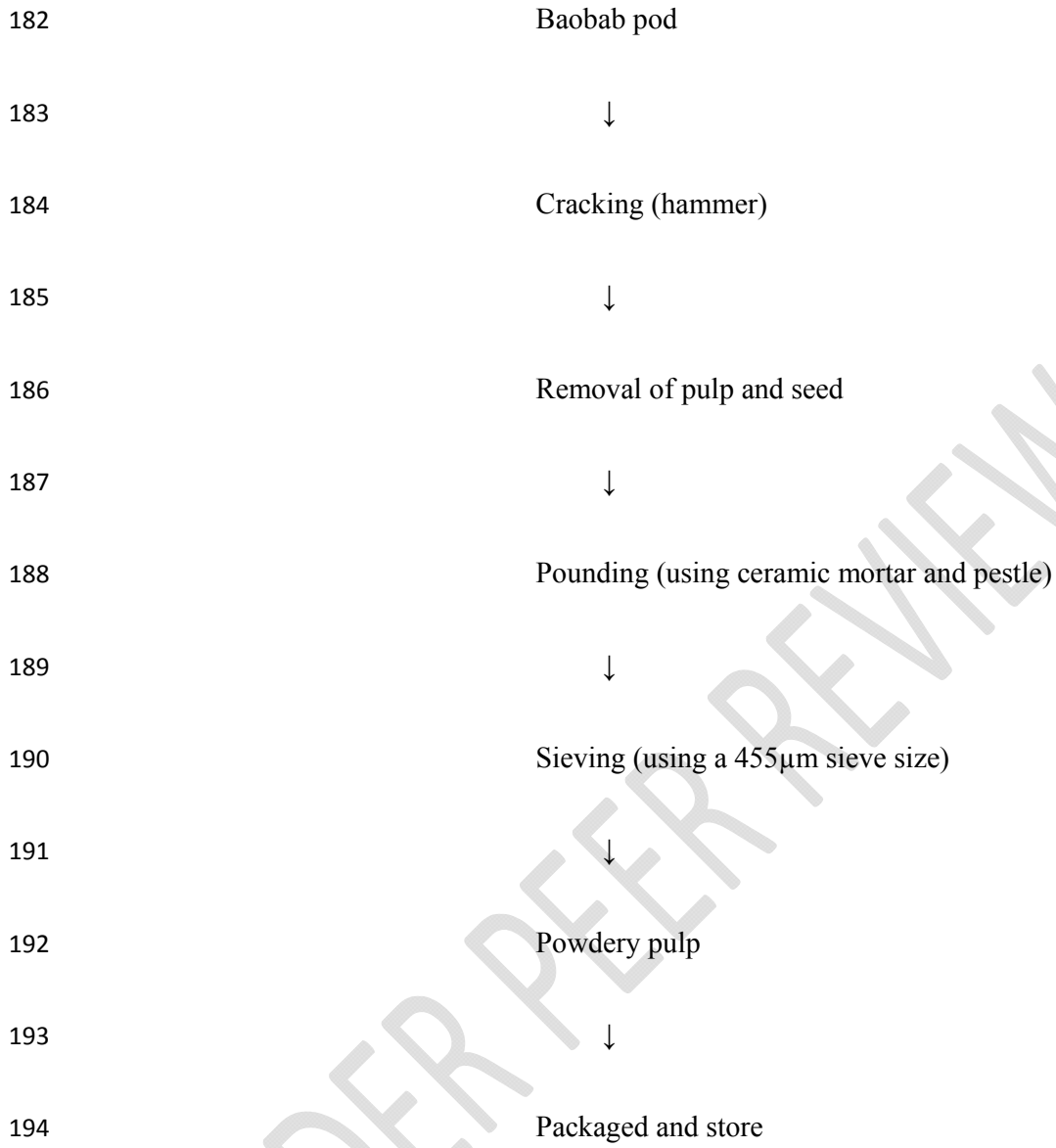
Packaged and store

180

Fig 2: Flow chart for the production of soy flour.

181

Source: Ihekoronye, 1999) with slight modification



195 Fig 3: Flow chart for the production of baobab fruit pulp powder.

196 Source: (Chadre, 2009) with slight modifications.

197

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199 .

200 **PROXIMATE ANALYSIS**

201 **Determination of Moisture Content**

202 Moisture content was determined by the air-oven method as described by AOAC (2005).
203 Two grams of the sample was weighed in duplicate into Petri dishes of know weight and
204 covered immediately. These were transferred into oven, uncovered and heated at $105^{\circ}\text{C} \pm 2$
205 for 3-5 hours. The samples were then removed from the oven and placed in the desiccator to
206 cool for 15 minutes before weighing. The process was repeated until constant weights were
207 recorded. The loss in weight from the original weight was reported as the moisture content.

$$208 \quad \% \text{ Moisture Content} = \frac{W_2 - W_3}{W_2 - W_1} \times 100 \quad (1)$$

209

210 **Determination of Crude Protein**

211 The Kjeldahl method was used for the determination of crude protein as described by AOAC
212 (2005). The samples (1.0g each) were first digested in Kjeldahl digesting system. The
213 digested samples were allowed to cool and then distilled into 2% boric acid solution
214 containing methyl orange indicator and diluted with water after the introduction of 40%
215 sodium hydroxide solution. The distilled samples were then titrated against 0.1 M HCL
216 solution. A blank titration was similarly carried out and the percentage content was estimated
217 as percentage Nitrogen $\times 6.25$ (1 ml of 0.1M HCL $\pm = 0.014$ g N)

$$218 \quad \%N = (b-a) \times 0.1N \text{ Hcl} \times 0.014 \times \text{dilution factor} \times 100 / \text{weight of sample} \quad (2)$$

$$219 \quad \% \text{ protein} = \% \text{ Nitrogen} \times 6.25 \quad (3)$$

220 **Determination of Crude Fat Content**

221 The Soxhlet solvent extraction method outlined in AOAC (2005) was used. Two gram
222 sample was weighed (A) into the extraction thimble and the thimble was plugged with cotton
223 wool. It was placed back in the Soxhlet apparatus fitted with a weighed flat bottom flask (B)
224 which was filled to about three quarter of its volume with petroleum ether of a boiling point
225 of $40\text{-}60^{\circ}\text{C}$. The extraction was carried for a period of 4-8 hours after which complete
226 extraction was made. The petroleum ether was removed by evaporation on the water bath and
227 the remaining portion in the flask was removed along with water by drying in the oven at 80
228 $^{\circ}\text{C}$ for 30 minutes and cooled in desiccators and weighed (C).

$$\% \text{ Fat Content} = \frac{W4 - W3}{W2 - W1} \times 100$$

229 where:

230 W1 = weight of oven dried thimble,

231 W2 = weight of sample used,

232 W3= weight of round bottom flask,

233 W4 = weight of round bottom flask with fat residue.

234 **Determination of Crude Fibre Content**

235 Fibre content was determined following the procedure outlined in AOAC (2005) method as
236 reported by Onwuka (2005) Two grams portions of the samples were extracted using
237 petroleum spirit (boiling point 40-60°C.) This was digested in 1 liter flask using 200ml
238 concentrated Sulphuric acid and filtered through the California buchner system .The insoluble
239 matter was washed with boiling water until it was free from the acid .The residue was then
240 back into the flask with 200ml of 0.313M NaOH. The flask content was brought to boil for
241 30 minutes. The flask was allowed to stand for 1 minute and filtered immediately through a
242 filtering cloth .The insoluble material was transferred into 100ml beaker by means of boiling
243 water, washed with 1% HCl and again with boiling water to free it from acid .The insoluble
244 material was finally washed with alcohol twice and three times with diethyl ether. The
245 resulting residue was transferred to a dish (previously weighed) with boiling water. The dish
246 containing the residue was dried for 2 hours, at 100°C, cooled in desiccators and weighed
247 (W1). The dried, cooled, and weighed residue was then transferred in a muffle furnace and
248 ignited at 600°C for 30 minutes, cooled and reweighed (W2). The percent crude fibre content
249 was calculated as follows.

250

$$251 \quad \% \text{ Crude Fibre} = \frac{W2 - W3}{W1} \times 100 \quad (5)$$

252 Where:

253 W1 = weight of sample used,

254 W2 = weight of crucible plus sample,

255 W3 = weight of sample crucible + ash.

256

257 **Determination of Ash**

258 The ash content of the sample was determined by the method described by AOAC (2005) a
259 silica dish was heated to 600⁰C, cooled in desiccators and weighed. Then 5g of the sample
260 was weighed into the silica dish and transferred to the furnace. The temperature of the
261 furnace was allowed to reach 525⁰C before placing the dish in it for 2 hrs. The temperature
262 was maintained until whitish grey colour was obtained indicating that all the organic matter
263 content of the sample had been destroyed. The dish was then brought out from the furnace
264 and placed in the desiccators, cooled and reweighed.

265

266
$$\% \text{ Ash Content} = \frac{W_2 - W_1}{\text{Weight of sample}} \times 100 \quad (6)$$

267 Where:

268 W₂ = weight of crucible + ash,

269 W₁ = weight of empty crucible.

270 **Determination of Carbohydrates**

271 Carbohydrate was determined by difference as reported by Ihekoronye and Ngoddy, (1985).

272
$$\% \text{ carbohydrate} = 100 - (\% \text{ moisture, protein, fibre, fat and ash}). \quad (7)$$

273 **FUNCTIONAL PROPERTIES OF SAMPLES**

274 **Determination of gelation capacity:**

275 The method described by Iwe *et al.* (2017) was used for the determination of the gelation
276 capacity. Suspensions of the samples in 5 ml of distilled water in test tubes were prepared
277 using 2 –20% (W/V) of the samples in test tubes. The sample test tubes were heated for 1
278 hour in a boiling water-bath followed by rapid cooling under running cold tap water. The test
279 tubes were further cooled for 2 hours at 40°C. Then, the gelation capacity was determined for
280 each sample as the least gelation concentration. That is, the concentration when the sample
281 from the inverted test tube will not slip

282 **Determination of Bulk Density**

283 The bulk density was determined as described by (Onwuka, 2005). A 10ml capacity
284 graduated measuring cylinder was weighed and 50g sample filled into it. The bottom of the

285 flask was tapped gently on the laboratory bench several times until there were no further
286 diminutions of the sample level after filling to 10ml mark.

$$287 \quad \text{Bulk Density (g/ml)} = \frac{\text{weight of sample}}{\text{volume of sample}} \quad (10)$$

288 **Determination of Swelling Index**

289 The method of Onwuka, (2005) was employed,. One gram of the flour samples was weighed
290 into 10ml graduated cylinder. Five (5ml) milliliters of distilled water was carefully added and
291 the volume occupied by the sample was recorded. The sample was allowed to stand
292 undisturbed in water for 1 hour and the volume occupies after swelling was recorded and
293 calculated as:

$$294 \quad \text{Swelling Index} = \frac{\text{vol.occupied by sample after swelling}}{\text{vol.occupied by sample after swelling}} \quad (11)$$

295 **Determination of Water Absorption Capacity**

296 The water absorption capacity of the flours was determine by the modified method of
297 Onwuka, (2005). One gram of sample was mixed with 10 mL distilled water and allow to
298 stand at ambient temperature (30 ± 2 °C) for 30 min, then centrifuged for 30 min at 3,000 rpm
299 or $2000 \times g$. Water absorption was examined as per cent water bound per gram flour.

300 **Determination of Oil Absorption Capacity**

301 The oil absorption capacity was also determined by the modified method of (Onwuka, 2005).
302 One gram of sample was mixed with 10 mL soybean oil (Sp. Gravity: 0.9092) and allow to
303 stand at ambient temperature (30 ± 2 °C) for 30 min, then centrifuged for 30 min at 300 rpm
304 or $2000 \times g$. Oil absorption was examined as percent water bound per gram flour.

305 **ENERGY VALUE**

306 This was calculated by multiplying the values of carbohydrate, fat and protein with the
307 Atwater Factor (4, 9, and 4) for carbohydrate, fat and protein respectively as described by
308 Onwuka, (2005).

309 **Sensory Evaluation**

310 Sensory evaluation based on the sensory attributes was conducted by using a standard 9-
311 points hedonic scales method (where 1 = dislike very much and 9 = like very much) as

312 described by Ihekoronye and Ngoddy, (1985). A total of 30 semi-trained panelists aged 18
313 years and above were involved in the evaluation of appearance, flavour, texture and overall
314 acceptability. The samples (100 g each) were coded randomly number using statistical
315 random Tables and served to the panellists with bottled water for rinsing their mouth after
316 every sample taste in a randomized order. The panellists were instructed to rate the attributes
317 indicating their degree of liking or disliking by putting a number as provided on the hedonic
318 scale according to their preference.

319

320 **Statistical Analyses**

321 All analyses were carried out in triplicate unless otherwise stated. Statistical significance was
322 established using one-way analysis of variance (ANOVA), and data were reported as the
323 mean standard deviation. Mean comparison and separation was done using Fisher's Least
324 Significant Difference test (LSD) at $p \leq 0.05$. Statistical analysis was carried out using the
325 SPSS 20 statistical package.

326

327

328

329 **DISCUSSION**

330 **Proximate Composition**

331 The proximate composition of sample A was significantly ($P < 0.05$) higher in protein content
332 (24.25%), fat content (16.65%) and Energy value (423.69 Kcal). According to Emmanuel *et al.*
333 *al.*, (2012), the addition of soybean flour to tiger-nut in the preparation of an infant diet
334 increases the protein, fat and energy values respectively. The Moisture content values for all
335 the samples tend to agree with the PAG (Protein Advisory Group – United Nations) which
336 reported moisture content of between 5-10% maximum. The range of moisture would have a
337 positive effect on the shelf life stability of the products (Bassey, 2004) and (Emmanuel *et al.*,
338 2012). The Ash content of the samples ranges from 2.59 – 2.87% with the highest value in
339 sample E (2.87%). The high Ash content of sample E could be due to the ratio of Millet Flour
340 and Baobab Fruit Pulp Powder in the sample since both are good sources of mineral
341 elements. Ash content of the samples was found to be less than the PAG standards which
342 reported 10% maximum ash content. The Protein content of the samples ranges from 9.80 –
343 24.25% with highest value in sample A (24.25%). These values are higher compared to PAG
344 standard (20%) respectively. This may be attributed to the protein content of soybean
345 addition (Emmanuel *et al.*, 2012). The fat content of the samples was found to range from
346 4.94 – 16.65% with sample A (16.65%) having the highest significance ($P < 0.05$) value than
347 others. This is as a result of the high soy (50%) flour content in the sample. Though, the fat
348 contents of sample A and B met the PAG standard which is 10% and for weaning foods.
349 Sample D and E with low Fat content could be as a result of low amount of soy flour addition
350 and increased baobab fruit pulp addition which may have caused some dilution. High Fat
351 content is very important in infant diet because it contain essential Fatty Acids (soy flour)
352 which promote good health. It is also a carrier of fat soluble vitamins (A, D, E and K) and
353 promoting the absorption (Emmanuel *et al.*, 2012). The Fibre content of the samples on the
354 other hand ranges from 4.62 – 11.65% with samples E (15.67%) having the highest
355 significant ($P < 0.05$) value. This could be due to increase in Baobab fruit pulp powder and
356 millet flour. An increase in the fibre content of weaning food has some beneficial effect on
357 the muscles of the large and small intestines. The values from the samples are higher than
358 those reported by PAG (5% Maximum). High fibre content was also reported to have adverse
359 effect on mineral element in the body (Emmanuel *et al.*, 2012) and (Bassey, 2004).

360 Carbohydrate content of the samples was found to range from 43.11-71.03% with sample E
361 having the highest significance ($P < 0.05$) value. The high values of carbohydrate could be as a
362 result of millet flour and possibly baobab fruit pulp. Carbohydrate is required in infant diet
363 for Energy during growth. Energy values of the samples was found to range from 367.78-
364 423.69 Kcal with sample A (423.69 Kcal) having the highest significance ($P < 0.05$) value.
365 The high Energy value of sample A is due to the high fat content of the sample. The Energy
366 value of the samples agrees with SON and PAG which reported 350-400Kcal respectively.
367 The Food and Agricultural Organisation reported that Home prepared weaning foods should
368 contain protein 15%, fat 11%, fibre 5% maximum, and for commercially prepared weaning
369 food for protein 15%, fat 6%, crude fibre 2% and moisture content 10% respectively.

370 **Functional Properties**

371 **Gelation concentration (GC)**

372 The least gelation concentration (LGC) which is defined as the lowest protein concentration
373 at which gel remained in the inverted tube was used as index of gelation capacity. The data
374 for LGC of different flours are given in Table 3. Composite (E) flours formed a gel at a
375 significantly higher concentration (10 g). Sample A and B flour formed gel quickly at very
376 lowest concentration (5 g). Wheat flours contain high protein and starch content and the
377 gelation capacity of flours is influenced by physical competition for water between protein
378 gelation and starch gelatinization (Kaushal *et al.* 2012). Suresh *et al.* (2015) reported that
379 protein gelation was significantly affected by exposed hydrophobicity and square of
380 sulfhydryls of proteins. As the percentage of incorporation of millet flour in wheat flour
381 (composite flour) increased, gelling properties decreased. The low gelation concentration of
382 A and B flour as composite flour may be added an asset for the formation of curd or as an
383 additive to other gel forming materials in food products. The variation in the gelling
384 properties may be ascribed to ratios of the different constituents such as protein,
385 carbohydrates and lipids in different flours, suggesting that interaction between such
386 components may also have a significant role in functional properties (Aremu *et al.* 2007). The
387 composite flours (E) would be useful in food system such as puddings, sauce and other foods
388 which require thickening and gelling (Suresh *et al.* 2015)

389 **Bulk density**

390 The bulk density (g/cm^3) of flour is the density measured without the influence of any
391 compression. The bulk densities of flours ranged from 0.69 g/cc to 0.71 g/cc. The highest
392 highest bulk density was observed A,B, C and D flour as shown in Table 3 and lowest was
393 sample E (0.69 g/cc). The present study revealed that bulk density depends on the particle
394 size and initial moisture content of flours. The obtained does not agree with those presented
395 by (Suresh *et al*, 2015), reported that Bulk density of composite flour increased with increase
396 in the incorporation of different flour. However, it is clear that decreased the proportion of
397 wheat flour increase the bulk density of composite flours. The high bulk density of flour
398 suggests their suitability for use in food preparations. On contrast, low bulk density would be
399 an advantage in the formulation of complementary foods (Suresh *et al*, 2015). Therefore, the
400 present study suggests that high bulk density of composite flour (A, B, C and D) suggests its
401 suitability to be used as thickener in food products and for use in food preparation since it
402 help to reduce paste thickness which is an important factor in convalescent and child feeding.

403 **Swelling capacity**

404 The swelling capacity of different flours ranged between 16.00 to 22.30 ml (Suresh *et al*,
405 2015). From Table 3, it is clear that lowest value of swelling capacity was observed in A
406 ($0.68 \pm 0.13 \text{ml}$) whereas the maximum in E ($1.04 \pm 0.13 \text{ml}$). The swelling capacity of flours
407 depends on size of particles, types of variety and types of processing methods and/or unit
408 operations. Suresh *et al*, (2015) reported that the flour of parboiled rice has more swelling
409 capacity as compared to raw rice. They also reported that the Swelling capacity of composite
410 flours increased with increase in the level of incorporation and decreased with level of wheat
411 flour addition. It is explicit that the swelling capacity of composite flours is highly affected
412 by the level of millet flour, because millet flour is rich in starch content.

413 **Water absorption capacity (WAC)**

414 The water absorption capacity for composite flours is given in Table 3. The WAC ranged
415 between 2.70 to 2.91 for all flours. The WAC was observed highest in C (2.91) and lowest in
416 D and E (2.70). The result suggests that addition of millet flour to wheat flour affected the
417 amount of water absorption. This could be due to molecular structure of millet starch which
418 inhibited water absorption, as could be seen from the lower values of WAC, with increase in

419 proportions of other flours to wheat flours. Similar observation was reported by Kaushal *et al.*
420 (2012). Suresh *et al.*, (2015) reported that lower WAC in some flours may be due to less
421 availability of polar amino acids in flours. The increase in WAC of blends after incorporating
422 millet flour may be due to increase in the amylose leaching and solubility and loss of starch
423 crystalline structure. High WAC of composite flours suggests that the flours can be used in
424 formulation of some foods such as sausage, dough and bakery products. The increase in the
425 WAC has always been associated with increase in the amylose leaching and solubility, and
426 loss of starch crystalline structure. The flour with high water absorption may have more
427 hydrophilic constituents such as polysaccharides. Protein has both hydrophilic and
428 hydrophobic nature and therefore they can interact with water in foods. The good WAC of
429 composite flour may prove useful in products where good viscosity is required such soups
430 and gravies. The observed variation in different flours may be due to different protein
431 concentration, their degree of interaction with water and conformational characteristics (Butt
432 and Batool, 2010).

433 **Oil absorption capacity (OAC)**

434 The composite flours (D and E) had highest OAC (2.72 and .44) and lowest for B (1.90). It is
435 clear that the OAC of composite flours increased with increase in the proportion of other
436 flours. The presence of high fat content in flours might have affected adversely the OAC of
437 the composite flours. The OAC was found to be insignificant to each other at $p \leq 0.05$ level of
438 significance. Therefore, the possible reason for increase in the OAC of composite flours after
439 incorporation of millet flour is the variations in the presence of non-polar side chain, which
440 might bind the hydrocarbon side chain of the oil among the flours. Similar findings were
441 observed by Kaushal *et al.* (2012). However, the flours in the present study are potentially
442 useful in structural interaction in food specially in flavor retention, improvement of
443 palatability and extension of shelf life particularly in bakery or meet products where fat
444 absorption is desired (Aremu *et al.* 2007). The major chemical component affecting OAC is
445 protein which is composed of both hydrophilic and hydrophobic parts. Non-polar amino acid
446 side chains which can form hydrophobic interaction with hydrocarbon chains of lipids
447 (Jitngarmkusol *et al.* 2008).

448

449

450 **Sensory Scores**

451 Table 4 shows the sensory scores of the samples tested. Appearance for sample A, B and C
452 was not significant ($P<0.05$) difference level but was significant ($P<0.05$) different level from
453 D and E. flavour shows that there are no significant ($P<0.05$) difference level in all the
454 samples tested. In terms of texture, there are no significant ($P<0.05$) difference level between
455 samples A, B and C and between samples B and C and also between sample C, D and D, E.
456 But there are significant ($P<0.05$) difference level between sample A and E, B and E and C
457 and D. the general Acceptability indicates that there are no significant difference ($P<0.05$)
458 between samples A, B, and C; samples B, C and D; samples C, D and E and between sample
459 D and E but there are significant difference ($P<0.05$) between sample A and E, B and E. The
460 sensory scores and general acceptability shows that sample A (7.66) was the most preferred
461 amongst all the tested sample followed by sample B (7.47) and C respectively.

462 **CONCLUSION**

463 The addition of baobab fruit pulp (BFP) to pearl millet and soybean flour, in turn increases
464 the fibre, ash and carbohydrate contents of the complementary foods. The functional
465 properties also improved with addition of baobab fruit pulp levels. This improvement could
466 be noticed in water absorption capacity, oil absorption capacity, bulk density and swelling
467 index. The sensory attributes indicates that the baobab fruit pulp samples competes very well
468 with the control (A) sample. However, sample A was most preferred by the panellist.

469 **Acknowledgement**

470 We wish to acknowledge all the Authors who articles, books etc we used.

471

Table 1: Blend Formulation of Pearl Millet, Soybean flour and Baobab Fruit Pulp (%)

Composite Flour

SAMPLES	MAIZE	SOYBEAN	BAOBAB FRUIT PULP
A	50	50	0
B	50	40	10
C	60	20	20
D	65	10	25
E	65	5	30

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473

UNDER PEER REVIEW

Table 2: Effect of Baobab Fruit Pulp Addition on the Proximate Composition a Complementary Food Samples.

SAMPLES	MOISTURE	PROTEIN	FAT	CARBOHYDRATE	FIBRE	ASH	ENERGY Kcal
A	10.98 ^c ±0.07	24.25 ^a ± 0.23	16.65 ^a ± 0.01	43.96 ^a ± 0.76	3.37 ^a ± 0.02	2.75 ^a ±0.00	423.69 ^a ±0.00
B	10.50 ^a ±0.02	20.38 ^a ± 0.18	13.90 ^b ± 0.08	43.11 ^a ± 0.34	7.68 ^b ±0.05	2.65 ^c ±0.03	379.06 ^b ±0.01
C	10.27 ^a ±0.06	14.58 ^b ± 0.30	8.84 ^c ± 0.00	62.00 ^b ±0.30	11.57 ^c ±0.08	2.68 ^a ±0.02	385.88 ^b ±0.03
D	10.73 ^a ±0.08	11.51 ^b ± 0.93	5.62 ^d ±0.04	67.91 ^b ±0.02	13.51 ^d ±0.06	2.59 ^b ±0.04	368.26 ^c ±0.00
E	10.09 ^b ±0.04	9.80 ^c ± 0.62	4.94 ^d ±0.02	71.03 ^c ±0.21	15.67 ^c ±0.05	2.87 ^a ±0.01	367.78 ^c ±0.02
LSD	0.08	0.06	0.02	0.01	0.02	0.09	0.08
PAG	5 - 10	20	10	-	5	10	350 - 400

Values are means of standard deviation. Values in the same column with different superscript are significantly (P,0.05) different

474 Key:

475 A = Millet 50%, soybean 50%

476 B = Millet 50%, soybean 40% and Baobab fruit pulp 10%

477 C = Millet 60%, soybean 20% and Baobab fruit pulp 20%

478 D = Millet 65%, soybean 10%and Baobab fruit pulp 25%

479 E = Millet 65%, soybean 5% and Baobab fruit pulp 30%

480 LSD = Least significant difference

481 PAG = Protein Advisory Group

482

Table 3: Effect of Baobab Fruit Pulp addition on The Functional Properties of a Complementary Food from Pearl Millet and Soy flour

SAMPLES	GELATION (%)	BULK DENSITY(g/ml)	Swelling Index (g/vol)	WAC	OAC
A	5.00±0.12	0.71±0.09	0.68±0.08	2.83±0.10	2.11±0.30
B	5.00±0.12	0.71±0.03	0.87±0.05	2.84±0.09	1.90±0.01
C	8.00±1.02	0.71±0.02	0.79±0.03	2.91±0.11	2.21±0.31
D	8.00±1.02	0.71±0.06	0.79±0.03	2.70±0.08	2.72±0.18
E	10.00±1.22	0.69±0.04	1.04±0.13	2.70±0.08	2.44±0.22

Means in the same column with different superscript are significantly ($p < 0.05$) different

484

485 **Key:**

486 A = Millet 50%, soybean 50%

487 B = Millet 50%, soybean 40% and Baobab fruit pulp 10%

488 C = Millet 60%, soybean 20% and Baobab fruit pulp 20%

489 D = Millet 65%, soybean 10% and Baobab fruit pulp 25%

490 E = Millet 65%, soybean 5% and Baobab fruit pulp 30%

Table 4: Effect of Baobab Fruit Pulp on The Sensory Attributes of a Complementary Food from Pearl Millet and Soy flour

SAMPLES	Appearance	Flavour	Texture	General Acceptability
A	7.26 ^a	6.60 ^a	6.53 ^a	7.66 ^a
B	7.20 ^a	6.40 ^a	6.33 ^a	7.47 ^a
C	7.13 ^a	6.00 ^a	6.07 ^b	7.20 ^b
D	6.53 ^a	5.73 ^c	5.40 ^d	6.73 ^b
E	5.80 ^c	5.27 ^d	4.67 ^d	5.33 ^c
LSD	0.974	1.390	1.334	1.086

Means in the same column with different superscript are significantly ($p < 0.05$) different

492 **Key:**

493 A = Millet 50%, soybean 50%

494 B = Millet 50%, soybean 40% and Baobab fruit pulp 10%

495 C = Millet 60%, soybean 20% and Baobab fruit pulp 20%

496 D = Millet 65%, soybean 10% and Baobab fruit pulp 25%

497 E = Millet 65%, soybean 5% and Baobab fruit pulp 30%

498 **COMPETING INTERESTS**
499 Authors have declared that no competing interests exist.
500

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