Nutritional quality of formulated complementary diet from defatted almond seed, yellow maize and quality protein maize flours

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Abstract

Traditional complementary foods are mainly starchy foods with limiting nutrient quality and can be fortified using protein rich crops like almond seed. This research thus aimed at investigating the nutritional quality of the formulated complementary diet from locally available almond seeds (Prunus amygdalus), high quality protein (QPM) and yellow maize after blending into flours. The proximate and amino acid compositions, in vitro protein qualities and functional properties of the blended flours were determined using standard methods. The in vivo studies involved feeding the weanling Wister albino rats with blended flours and commercial Cerelac (control), followed by hematological and histopathological determinations, while sensory attributes were evaluated by the semi-trained panelists. The protein contents of the flour blends (24–28%) were significantly (p < 0.05) better with adequate indispensable amino acids and improved functionalities than commercial cerelac (23%). Comparatively, the dried germinated QPM (DGQPM) has significant (p < 0.05) higher biological value (~37%) than fermented high QPM (FHQPM) (~30%), thereby indicating that the germination process improved protein quality of the diets. Besides, the in vivo data showed a positive effect of germination process as the rats fed with DGQPM has low white blood cells (30 × 10^2) compared to FHQPM (42 × 10^2) and cerelac (51 × 10^2). However, the fermentation process improved the packed cell volume of rats fed with FHQPM (49%) when compared to DGQPM and cerelac (47%). The formulated diets have no negative effects on the protein content (45.19–51.88 mg N/g) and weight (0.25–1.36 g) of the internal organs (liver, kidney and tissue) of the animals when compared to cerelac (53.72–55.04 mg N/g; 0.25–1.98 g), respectively. The panelists generally accepted all the formulated diets, hence encouraging their utilization in the global preparation of complimentary foods for young children to meet their nutritional needs and adding value to the locally produced underutilized almond seeds.

Keywords: Quality protein, Yellow maize, Almond seed, Nutrient composition, Complementary

Introduction

An adequate nutrition within the first 1000 days has been reported to be essential for healthy growth and development in children for their full potential (Motuma et al. 2016). The consequences of poor nutrition include illness such as common childhood diarrhea, global malnutrition (Onyango et al. 2014), stunting (a state of an adult being shorter than potential height), and micronutrient deficiencies (Motuma et al. 2016). Therefore, the introduction of protein and energy-rich complementary foods to the children is very critical at this stage in order to avert the challenges of good nutrition faced by the children of 6–23 months in most developing countries (Udoh and Amodu 2016). Besides, highly nutritious diets
with rich amino acid contents, such as arginine, tyro-
sine/phenylalanine and valine/leucine, have the capacity to
fight against the chronic cardiovascular diseases. For in-
stance, arginine, aromatic and branched-chain
amino acids have been extensively reported for their
health beneficial effects such as, antioxidant (Malomo
et al. 2020), antihypertensive (Poggiogalle et al. 2019)
and adequate combating agents towards protein-energy
malnutrition commonly associated with growing chil-
dren (Arise et al. 2020).

Over a very long period of time in the developing
countries, cereal based complementary food often fail to
meet the nutritional need of younger children due to
poor/low nutrient contents (Ijarotimi and Keshinro
2012a). Such cereals like sorghum, rice and maize con-
stituted about 85% of total global cereals utilized in
preparation of local complementary foods (Sofi et al.
2009) but with implications of having low protein con-
tents (Tufa et al. 2016). However, some average-income
earners nowadays prepare their complementary foods
from cereals mixed with roasted soybeans and crayfish
flours (Alamu et al. 2018). This practice might be very
expensive for most low-income earners or the rural
dwellers but who could find the locally available and
underutilized almond seeds useful for their own prepara-
ration.

Almond seeds contain about 21.2% protein with 30 g
serving of almonds providing 6.3 g of protein (Ahren
et al. 2005; United States Department of Agriculture,
USDA 2018) with lots of phytosterols and ranked number
two after cashew nuts in tree nut (Shakerardekani
et al. 2013). Studies had shown that besides high nutriti-
ve values, almonds provide varied healthful effects like
cardiometabolic, modulation of blood glucose fluctua-
tions, reduction in postprandial plasma lipids and free
radicals scavengers (Tan and Mattes 2013; Esquius et al.
2020; Palacios et al. 2020; Semidei et al. 2020).

Quality protein maize (QPM) is a bio-fortified maize
with higher amount (46 and 66% higher than common
maize) of lysine and tryptophan (the first and second
limiting amino acids in maize), respectively (Ikuenjola
and Ogunba 2018) but lower amount of leu-
cine and isoleucine (Fagbemi et al. 2018). These two
amino acids (lysine and tryptophan) are found indispens-
able for healthy growth and developments of infants and
young children (Ijarotimi and Keshinro 2012a). Such a
balanced combination of amino acids in the endosperm
resulted into its higher biological value ensuring more
availability of protein to human being than common
maize (Ikuenjola and Ogunba 2018). Although, the total
amount of protein in QPM is not actually increased but
the protein is enhanced so that it delivered a higher
benefit when consumed by human beings (Fagbemi et al.
2018). QPM has previously been used to produce
breakfast and weaning food blends (Abiose and Ikujenlola
2014; Ikuenjola and Ogunba 2018; Krishna-Motukuri 2019;
Gemede 2020; Parsons et al. 2020).

The nutritional problems arising from wide utilization
of common maize, sorghum and guinea corn with poor/
low protein contents, limiting lysine and tryptophan as
well as expensive purchasing cost of high protein
sources (Ikuenjola and Ogunba 2018) had paved way
for the utilization of almond seeds and QPM. Therefore,
there is need to explore the nutritional qualities of these
crops in the production of complementary food for chil-
dren due to their nutrient composition and accessibility
to many low-income homes. Besides, the germination
of QPM has the advantage of breaking down the complex
sugars and other matrix of the nutrients available within
the QPM, to make more protein and amino acids avail-
able (Mares et al. 2018), while fermentation caused the
reduction of these organic portions through the increase
in the metabolic activities of the microbes (Olagunju
and Ifesan 2013).

Although, significant information abound globally on
the use of composite flour blends from different cereal
and legume crops but there exist scanty information on
the utilization of almond seed and QPM blends as com-
plementary food for children. Hence, the aim of this
study was to formulate and assess the nutritional quality
of almond seed, QPM and yellow maize flour blends as
ingredients for complementary food.

Materials and methods
Materials
The almond fruits (Prunus amygdalus) were collected
fresh from the Federal Polytechnic, Ado-Ekiti, Ekiti
State, Nigeria. High quality protein and yellow maize
were purchased from Institute of Agricultural Research
and Training, Moor Plantation, Apata Ganga, Ibadan,
Oyo State, Nigeria. The weanling Wister albino rats
(150–185 g) were purchased from Department of Bio-
chemistry, Obafemi Awolowo University, Ile-Ife, Osun
State, Nigeria. N-hexane, other chemicals and reagents
used were purchased from Rovet Stores, Akure, Ondo
State, Nigeria. The Cerelac (a commercial product) and
basal diet (both control samples) were obtained from
NAO supermarket, Akure, Ondo State, Nigeria.

Processing of de-fatted almond seed to flour
The almond seeds were defatted and processed into
flour using the method previously described by Tan and
Mannes (2013). The edible portion of the fruit was manu-
ally removed leaving the stony shell containing the seed.
The stony shell was carefully cracked using harmer to
remove the nut. The almond (nuts) were sorted and
dried in hot air oven at 60°C until dried, and then
milled, using a Philips laboratory blender (Philips
HR2811 model, Amsterdam, Netherland). The resulting meal was defatted using n-hexane in a soxhlet extraction apparatus as described by AOAC (2005). The defatted sample was air-dried in a fume hood at room temperature to drive off the n-hexane completely, the flour was blended by using a blender and thereafter sieved and preserved in tightly closed plastic container pending analysis.

**Processing of maize into fermented and germinated maize flours**

The fermented maize flour was prepared according to the method previously described by Ijarotimi and Keshinro (2012b) with slight modification. The maize grains were cleaned, sorted, and steeped in water for 72 h to naturally ferment. The steep water was decanted and the maize was washed in clean water after which they were wet-milled using hammer mill. After milling, the slurry was sieved with cheese-cloth and the coarse particle was discarded and the slurry was settled for 3 h. The washed water was decanted, the sediment was transferred into a clean cheese-cloth and pressed to remove excess water by using locally manufactured hydraulic press. The sediment was dried in an air oven at 50 °C for 48 h and re-milled into fine flour and stored in air-tight plastic container in a cool dry place for further analysis and formulation.

The germinated maize flour was obtained according to the method previously described by Ijarotimi and Keshinro (2012a). Briefly, the maize grains were soaked in water for 2 h to achieve hydration, after which it was then left at room temperature (25 °C) for 48 h. This was thereafter rinsed and spread on a jute bag for germination to take place while the seeds were closely monitored, frequently watered and separated so as to prevent mold growth. After 48 h the seeds were rinsed and the germination process was terminated by exposing the seeds to a temperature of 40 °C for 5 h. The germinated seeds were milled into flour using hammer mill and then sieved through a 0.4 mm mesh size, packaged in high-density polyethylene bag and stored in a refrigerator at 4 °C till further use.

**Formulation of the complementary diets**

The complementary diets were made from blends of 70% maize (QPM or yellow maize) and 30% defatted almond flour as previously described by Ijarotimi and Keshinro (2012a). The mixes obtained were thoroughly blended together using Philips laboratory blender (Philips HR2811 model, Amsterdam, Netherland). After blending, the blend mixes were packed and sealed in high-density polythene bags until required while cerelac was used as a control.

**Proximate composition analysis**

The proximate analysis (moisture, protein, fat, ash and crude fiber) of the diets was determined according to the methods of Association of Official Analytical Chemists (AOAC 2005) while the carbohydrate content was calculated by difference. The crude protein and fat contents were obtained by micro-Kjedahl (N × 6.25) and soxhlet procedures, respectively.

**Amino acid composition analysis**

Amino acid content was determined using Pico-Tag method as previously described by Bidlingmeyer et al. (1984). The cysteine and methionine (sulphur-containing amino acids) were determined after performic acidoxidation as described by Gehrke et al. (1985) and tryptophan was determined after alkaline hydrolysis as described by Landry and Delhaye (1992).

Briefly, sample was hydrolysed, evaporated in a rotary evaporator and loaded into Technicon Sequential Multi-Sample Amino Acid Analyser (TSM-1) (Technicon Instruments Corporation, New York, USA). 10 μL of each hydrolysate was dispensed into the cartridge of the analyser followed by 76h of free acidic, neutral and basic amines analysis. Norleucine was employed as the internal standard. Ten microliter (10 μL) of the standard solution mixture of the amino acid was also loaded into the analyser. Values of both the standard and samples was recorded and printed out as chromatogram peaks by the chart recorder.

**Calculation from the peaks:** The net height of each peak produced on the chromatogram (each representing amino acid) was measured. The half-height of each peak was located and the width of the peak at half-height will accurately be measured. Approximate area of each peak was then obtained by multiplying the height with the width of the half height.

**Functional property analysis**

The functional properties were determined according to the procedures previously described by Arise et al. (2018). The bulk density (BD) was determined when 10 g of sample was placed in a graduated cylinder (50 ml) and packed by gently tapping the cylinder on the bench top (10 times) to form a reasonable height. The volume of sample was recorded while BD was expressed as grams material per milliliter. For the determination of swelling power, briefly, 2 g of the sample was weighed and poured into a measuring cylinder and the initial value was noted, 25 ml of distilled water was added, the solution was vigorously shaken and left for 30 mins, 1 and 2 h. The final reading of the swelled sample was taken. The percentage of the swelling index was calculated as:
Swelling power ($\%$) = \( \frac{\text{Initial reading}}{\text{Final reading}} \times 100 \)

The water/oil absorption capacity was determined when 1 g of each sample was dispersed in 10 mL of distilled water/oil in a 50-mL preweighed centrifuge tube. The dispersion was vortexed for 1 min, allowed to stand for 30 min and centrifuged at 4000 x g for 30 min at room temperature. The supernatant was decanted, excess water/oil in the upper phase was drained for 15 min, and the tube containing the residue was weighed again to determine the amount of water or oil retained per gram of the sample.

\[ \text{WAC or OAC} \% = \frac{\text{Volume of water/oil used} - \text{volume of free water/oil}}{\text{Weight of sample used}} \times 100 \]

The emulsifying activity was determined by the method of Wang and Kinsella (1976), with modifications. To verify the emulsifying activity, 20 ml of maize oil was added to 20 ml of suspension protein and subsequently stirred with domestic mixer (Kenwood Appliances, Woking, Surrey, England) for 1 min. The emulsion formed was divided into tubes and centrifuged at 4000 rpm for 6 min. The emulsifying activity was determined as the percentage of the emulsified layer that remained after centrifugation, and the percentage calculation performed using:

\[ \text{WAC or OAC} \% = \frac{\text{Mass of the emulsified layer}}{\text{Weight of sample used}} \times 100 \]

The least gelation concentration was done using test tubes containing suspensions of 2, 4, 6, 8 up to 20% (w/v) flour in 5 ml distilled, heated for 1 h in boiling water, followed by cooling in ice and further cooling for 2 h at 4°C. The viscosity was done using capillary tubes and the time it takes for a volume of liquid to pass through the length of the tube.

In vivo studies

Ethical clearance

The study protocol was approved and ethic clearance given by the Ethical Committee for Laboratory Animals of School of Agriculture and Agricultural Technology, Federal University of Technology, Akure, Nigeria (FUTA/SAAT/2016/033).

Experimental animals

A total of thirty (30) weanling Wister albino rats purchased from Department of Biochemistry, Obafemi Awolowo University, Ile-Ife, Osun State, Nigeria were used for the experiment according to the methods of Odebode et al. (2017). Before the arrival of the rats, the rat house and cages were properly cleaned and disinfected using disinfectant. The cages were properly arranged and fitted with feeders and drinkers that can comfortably drop water when nibbled by the rats. The feeders were firmly placed in positions to eliminate feed spillage. The rats were weaned at about 21–25 days, weighing 30–65 g. They were thereafter divided into six groups of five rats each on the basis of initial weight. The rats were housed individually in stainless steel metabolic cages with facilities for separate collection of feaces and urine.

Experimental diet

Four different experimental diets were developed using maize and almond seed. Fermented yellow maize (FYM) and fermented high quality protein maize (FHQPM) grains were processed into fermented maize flour. Dried germinated high quality protein maize (DGHQPM) and dried high quality protein maize (DHQPM) grains equally processed to maize flour. All the processed maize flours were mixed with almond seed flour at ratio of 70 and 30%, respectively. Basal diet (nitrogen-free diet) was formulated according to Fanimo (Fanimo, A: Substitution of Soyabean and animal by-products for fish meal in pig rations Ph.D Thesis Department of Animal Science. Ibadan, Oyo State: University of Ibadan. Unpublished), such that there was no nitrogen furnished by any of the ingredients. One group of five rats was given nitrogen-free diet (negative control); another four groups of five rats were fed on the remaining four diets containing the test ingredients while the last group was given cereals as positive control.

Slaughtering and collection of blood samples

At the 28th day of the experiment, all the rats were starved for about 3 h and weighed. Each rat was anaesthetized with chloroform inside a dessicator before slaughtering. Blood was collected into bijour bottle containing a speck of dried ethylene diamine tetrachloro-acetic acid (EDTA) powder as anti-coagulant. The bottles were immediately capped and the content mixed gently for about 1 min by repeated inversion thereafter used for various haematological studies.

Measuring of rat organs (Histopathological)

The organs: heart, kidney and liver were weighed and expressed in g for each rat.

Haematological studies

From the blood collected, the packed cell volume (PCV) was estimated by spinning about 75 ml of each sample in heparinized capillary tubes in a hematocrit micro-centrifuge for 5 min while the total white blood cells (WBC) were determined as described by Lamb (1981). The lymphocytes, neutrophils, monocytes, basophils, platelets
and eosinophils were also determined as described by Lamb (1981).

Sensory evaluation
The formulated diets and the control sample were subjected to sensory evaluation as described by Olapade and Aworh (2012). The samples were reconstituted, coded and served warm to ten (10) semi-trained panelist drawn from caregivers (Nursing mothers) attending Child Welfare Clinic in a Comprehensive Health Centre, Ado-Ekiti, Ekiti State, Nigeria. Each panelist was given coded samples to rate for appearance, aroma, taste, consistency and overall acceptability using 9-point Hedonic scale from dislike extremely (1) to like extremely (9).

Statistical analysis
Triplicate replications were used to obtain mean values and standard deviations. Statistical analysis was performed with SAS (Statistical Analysis Software 9.1) using one-way ANOVA. Duncan’s multiple-range test was carried out to compare the mean values for samples with significant differences taken at $p < 0.05$.

Results and discussion
Proximate composition of formulated flour blends
The proximate composition is important in determining the quality of food raw materials and often the basis for establishing the nutritional value and consumers’ overall acceptance of the product (Mashood and Rizwana 2010). Hence, the proximate compositions of yellow maize, high quality protein maize (QPM) and almond flours blends are presented in Table 1. The blends were all dried to a minimum level (6–7%) found in the range of the FAO recommended level (<10%) for most dried food samples (National Research Council, NRC 2009), while the remaining compositions were calculated based on the dry weights of the blends. The crude protein content ranged from 24.28 to 27.64% with the dried germinated QPM (DGQPM) blend having significantly ($p < 0.05$) highest value (27.64%) when compared to 26.77, 26.62 and 23.29% for dried QPM (DQPM), fermented yellow maize (FYM) and cerelac (commercial control) samples, respectively. The requirements for the maintenance of body protein equilibrium and the optimum pattern of individual indispensable amino acids deviated literally between the ages of 6 and 24 months (Reeds and Garlick 2003), hence the need to absorb quality and concise amount of protein at the age of <6 months. The high protein content of DGQPM (~28%) could be attributed to malting of the maize, which encouraged the crop’s improved protein as previously reported (Ijarotimi and Keshinro 2012a). However, the current protein contents were above the recommended protein level (20%) reported for formulated foods (Akinola et al. 2014) and significantly ($p < 0.05$) higher when compared with 26.87% reported for fermented-popcorn-African locust bean-Bambara complementary foods (Ijarotimi and Keshinro 2012b). The crude fat, total ash, crude fibre and carbohydrate contents of the formulated blends (4.69–9.82, 0.55–2.16%, 0.94–0.97, 62.77–64.67%, respectively) presented in Table 1 were all within the ranges (10–70%) recommended for infants and young children (National Research Council, NRC 2009). The implication of this data is that these blends would pose no undesirable bulk density, low caloric density, irritation of the gut mucosa and adverse effects on the efficiency of absorption of various significant nutrients. All these undesirable factors had previously been associated with Ogi, the common traditional complementary foods (Onabanjo et al. 2008).

Amino acid composition of formulated complementary flour blends
The amino acid content of complementary foods is a particularly relevant issue to infant feeding in developing countries with continual protein-energy malnutrition challenges (Onabanjo et al. 2008). This might be due to the poor feeding practices and low quality protein commonly associated with plant-based diets (Badamosi et al. 1995). The amino acid compositions of the formulated complementary flour blends are presented in Table 2. The data showed that the dried germinated QPM (DGQPM) and fermented high QPM (FHQPM) had the highest (5.81 g/100 g) and least (3.76 g/100 g) tryptophan contents, respectively. The current result (3.76–5.81 g/100 g) is higher than 1.95, 1.31 and 1.25 g/100 g reported

### Table 1 Proximate compositions (d.w.) of blends of maize and almond flours (%)

| Samples      | Moisture content | Total ash | Crude fat | Crude fibre | Crude protein | Carbohydrate |
|--------------|------------------|-----------|-----------|-------------|---------------|--------------|
| DQPM         | 6.74 ± 0.02a     | 2.01 ± 0.02a | 7.27 ± 0.04a | 0.97 ± 0.02a | 26.77 ± 0.03a | 62.99 ± 0.13a |
| DGQPM        | 6.15 ± 0.02a     | 2.07 ± 0.04a | 4.69 ± 0.40a | 0.94 ± 0.02a | 27.64 ± 0.04a | 64.67 ± 0.48a |
| FYM          | 6.54 ± 0.04c     | 0.55 ± 0.02c | 8.00 ± 0.04b | 0.96 ± 0.01a | 26.62 ± 0.03a | 63.87 ± 0.10c |
| FHQPM        | 7.24 ± 0.03a     | 2.16 ± 0.01b | 9.82 ± 0.20a | 0.97 ± 0.03a | 24.28 ± 0.02a | 62.77 ± 0.05a |
| CERC         | 3.32 ± 0.02a     | 3.38 ± 0.03a | 4.49 ± 0.02d | 0.62 ± 0.01b | 23.29 ± 0.02a | 68.20 ± 0.09a |

DQPM Dried Quality Protein Maize, DGQPM Dried Germinated Quality Protein Maize, FYM Fermented Yellow Maize, FHQPM Fermented High Quality Protein Maize, CERC Cerelac. Mean values with different superscript on the same column are significant ($p < 0.05$)
for commercial cerelac (control), fermented popcorn—wonderful kola seed flour blend (Ijarotimi et al. 2015) and USDA Recommended Daily Allowance (United States Department of Agriculture, USDA 2018). The higher tryptophan content obtained for the formulated-QPM blends when compared to other products from common maize might be due to the contribution of reported higher tryptophan content of QPM than common maize (Ikujenlola and Ogunba 2018). However, FHQPM has the highest glutamic (9.06 g/100 g) and aspartic (6.54 g/100 g) acid when compared to dried quality protein maize (DQPM) (2.65; 2.65 g/100 g) and cerelac (3.29; 2.55 g/100 g), respectively. The fermentation process might have positively affected the high proportions of these acidic amino acids in FHQPM. Meanwhile, the DQPM has the highest total indispensable amino acid (35.01 g/100 g) while FHQPM has the least (27.28 g/100 g) as shown in Table 2. Although, all the indispensable amino acids were present in the amino acid profile but quite a number did not meet the FAO and WHO (1991) referenced values. A lower score for any of the indispensable acids designated the limiting characteristics of the amino acids and gave an indication of the protein quality of such plant-based food materials when compared with food materials produced from animal sources (Ogunlade et al. 2005).

### Functional properties of the formulated flour blends

The functional property of food material is an important determinant in the application and use of such food material for various food productions (Alawode et al. 2017). Table 3 showed the functional properties of the formulated complementary foods in comparison with the control. The least gelation concentration (LGC) of the samples ranged from 2 to 6% with the least LGC (2%) obtained for dried quality protein maize (DQPM), fermented yellow maize (FYM) and fermented high quality protein maize (FHQPM), respectively. This implied that the DGQPM might possess reduced viscosity, plasticity and elasticity, thus forming a low dietary bulk, which is highly favourable for a good complementary diet (Omueti et al. 2009). Studies have shown that foods that formed gels at low concentrations

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Table 2 Amino acid compositions (g/100 g) of blend of maize and almond seed flours

| Amino acids  | Samples    | DQPM  | DGQPM | FYM  | FHQPM | CERC | *RDA |
|-------------|------------|-------|-------|------|-------|------|------|
|             |            | 3.09 ± 0.02<sup>c</sup> | 3.78 ± 0.32<sup>b</sup> | 3.74 ± 0.02<sup>b</sup> | 3.01 ± 0.02<sup>c</sup> | 4.09 ± 0.02<sup>a</sup> | – |
| Alanine     |            | 1.97 ± 0.06<sup>b</sup> | 5.61 ± 2.86<sup>a</sup> | 2.16 ± 0.02<sup>b</sup> | 6.54 ± 0.02<sup>a</sup> | 2.55 ± 0.02<sup>b</sup> | – |
| Aspartic acid|            | 2.65 ± 0.02<sup>a</sup> | 6.88 ± 0.02<sup>a</sup> | 3.05 ± 0.02<sup>d</sup> | 9.06 ± 0.02<sup>c</sup> | 3.29 ± 0.02<sup>f</sup> | – |
| Glutamic acid|            | 2.89 ± 0.02<sup>d</sup> | 3.11 ± 0.02<sup>b</sup> | 3.35 ± 0.02<sup>a</sup> | 2.66 ± 0.09<sup>d</sup> | 3.28 ± 0.02<sup>d</sup> | – |
| Glycine     |            | 2.89 ± 0.02<sup>d</sup> | 2.97 ± 0.02<sup>c</sup> | 3.35 ± 0.02<sup>a</sup> | 2.12 ± 0.02<sup>d</sup> | 3.28 ± 0.02<sup>b</sup> | – |
| Proline     |            | 8.50 ± 0.02<sup>b</sup> | 2.40 ± 0.02<sup>d</sup> | 9.31 ± 0.02<sup>a</sup> | 1.81 ± 0.02<sup>e</sup> | 7.30 ± 0.02<sup>e</sup> | – |
| Serine      |            | 3.83 ± 0.02<sup>c</sup> | 2.16 ± 0.02<sup>c</sup> | 4.08 ± 0.02<sup>a</sup> | 1.94 ± 0.02<sup>d</sup> | 4.08 ± 0.02<sup>a</sup> | 1.9 |
| Histidine   |            | 2.32 ± 0.02<sup>d</sup> | 2.86 ± 0.02<sup>d</sup> | 3.01 ± 0.02<sup>b</sup> | 2.20 ± 0.02<sup>d</sup> | 3.11 ± 0.02<sup>a</sup> | 2.8 |
| Isoleucine  |            | 6.51 ± 0.02<sup>c</sup> | 7.25 ± 0.02<sup>b</sup> | 2.89 ± 0.02<sup>e</sup> | 5.66 ± 0.02<sup>d</sup> | 8.01 ± 0.02<sup>a</sup> | 6.6 |
| Leucine     |            | 2.85 ± 0.02<sup>b</sup> | 3.17 ± 0.02<sup>a</sup> | 3.06 ± 0.02<sup>a</sup> | 2.74 ± 0.05<sup>d</sup> | 3.04 ± 0.05<sup>a</sup> | 5.8 |
| Lysine      |            | 1.22 ± 0.02<sup>d</sup> | 1.59 ± 0.02<sup>d</sup> | 1.30 ± 0.02<sup>c</sup> | 0.96 ± 0.02<sup>d</sup> | 2.08 ± 0.02<sup>a</sup> | 2.2 |
| Methionine  |            | 3.38 ± 0.02<sup>b</sup> | 4.06 ± 0.02<sup>b</sup> | 3.97 ± 0.02<sup>c</sup> | 3.13 ± 0.02<sup>d</sup> | 4.23 ± 0.02<sup>e</sup> | 2.8 |
| Phenylalanine|            | 2.05 ± 0.02<sup>d</sup> | 2.33 ± 0.02<sup>b</sup> | 2.19 ± 0.02<sup>c</sup> | 1.80 ± 0.02<sup>d</sup> | 2.27 ± 0.01<sup>b</sup> | 3.4 |
| Threonine   |            | 6.86 ± 0.02<sup>b</sup> | 4.34 ± 0.02<sup>c</sup> | 6.98 ± 0.07<sup>e</sup> | 3.66 ± 0.02<sup>d</sup> | 2.55 ± 0.02<sup>c</sup> | 2 |
| Arginine    |            | 0.82 ± 0.02<sup>d</sup> | 0.99 ± 0.02<sup>b</sup> | 0.93 ± 0.02<sup>c</sup> | 0.53 ± 0.02<sup>b</sup> | 1.39 ± 0.01<sup>f</sup> | – |
| Cystine     |            | 2.25 ± 0.02<sup>e</sup> | 2.42 ± 0.02<sup>b</sup> | 2.25 ± 0.02<sup>c</sup> | 1.93 ± 0.02<sup>d</sup> | 2.58 ± 0.02<sup>e</sup> | – |
| Tyrosine    |            | 4.72 ± 0.01<sup>b</sup> | 5.81 ± 0.02<sup>a</sup> | 4.79 ± 0.01<sup>c</sup> | 3.76 ± 0.02<sup>d</sup> | 1.95 ± 0.02<sup>d</sup> | 1.25 |
| TSAA (Meth + Cys)| 2.04 | 2.58 | 2.23 | 0.94 | 3.47 | 2.20 |
| AAA (Phe + Tyr + Trp)| 6.35 | 7.29 | 6.98 | 5.85 | 7.76 | 2.80 |
| TIAA       |            | 35.73 | 35.09 | 34.73 | 28.07 | 36.94 | 33.90 |

*DQPM Dried Quality Protein Maize, DGQPM Dried Germinated Quality Protein Maize, FYM Fermented Yellow maize, FHQPM Fermented High Quality Protein Maize, CERC Cerelac, TSAA Total Sulphur Amino Acid, AAA Aromatic Amino Acid, TIAA Total Indispensable Amino Acid, *RDA Recommended Daily Allowance (United States Department of Agriculture, USDA 2018). Values with different superscript on the same row are significant (p < 0.05)*
were not ideal for infant diet, because of the high dilution rate in an attempt to improve digestibility-volume relation (Ezeji and Ojimelukwe 1993; Obatolu and Cole 2000).

Water absorption capacity (WAC) is an index of the maximum amount of water that a food product would absorb and retain (Marero et al. 1988). It is also the ability of a product to associate with water under a limiting water condition. WAC is desirable for food systems to improve yield and consistency as well as given body to the food (Osun-dahunsi et al. 2003). The WAC of the formulated diets ranged from 150 to 295 g/g with DQPM and FYM having the highest values, respectively, which might be due to their reported protein contents (Table 1). However, the lowest WAC of DGQPM might be as a result of the microbial activities of the flour blend but with a resultant extended shelf-live (Giami and Bekeham 1992).

Oil absorption capacity (OAC) of the flour blend is the required protein (Jachmanian et al. 1995). Moreover, the broken down and prevented the bioavailability of the required protein (Jachmanian et al. 1995). Moreover, the

Table 3 Functional properties of blends of maize and almond seed flours

| Functional properties               | Samples       | DQPM | DGQPM | FYM  | FHQPM | CERC |
|-------------------------------------|---------------|------|-------|------|-------|------|
| Least gelation (%)                  |               | 2.00 | 6.00  | 2.00 | 2.00  | 4.00 |
| Water absorption capacity (g/g)     |               | 295.00 ± 5.00b | 150.00 ± 0.00d | 295.00 ± 5.00b | 240.00 ± 10.00d | 350.00 ± 10.00d |
| Oil absorption capacity (g/g)       |               | 214.66 ± 4.50b | 264.66 ± 9.50a | 173.00 ± 9.00g | 182.00 ± 0.00d | 205.66 ± 4.50d |
| Bulk density (g/ml)                 |               | 0.80 ± 0.00  | 0.64 ± 0.05b  | 0.54 ± 0.01c  | 0.66 ± 0.00b  | 0.77 ± 0.02a  |
| Emulsion capacity                   |               | 52.50 ± 0.50a | 51.00 ± 1.00b  | 50.50 ± 1.00b  | 51.50 ± 1.50b  | 51.00 ± 1.00b  |
| Swelling power (g/g)                |               | 3.10 ± 0.10a | 2.50 ± 0.10c  | 2.85 ± 0.10b  | 2.58 ± 0.08c  | 3.02 ± 0.07a  |
| Viscosity (cPas)                    |               | 1.50 ± 3.00a | 1.70 ± 2.00a  | 1.70 ± 8.00a  | 1.70 ± 8.00a  | 1.50 ± 3.00a  |

DQPM Dried Quality Protein Maize, DGQPM Dried Germinated Quality Protein Maize, FYM Fermented Yellow Maize, FHQPM Fermented High Quality Protein Maize, CERC Cerelac. Values with different superscript on the same row are significant (p < 0.05)

The swelling power (SP) capacity is an important factor used to determine the amount of water that food samples would absorb as well as the degree of swelling within a given time (Ijarotimi and Keshinro 2012a). The SP ranged from 2.50 to 3.10 with no significant difference (p > 0.05) between the control and DQPM diets, whereas, DQPM and DGQPM blends have the highest and least SP, respectively. This implied that the DQPM with the highest SP when compared to the other complementary diets (DGQPM, FYM and FHQPM) would produce a thick viscous gruel.

Nutritional quality of formulated flour blends

The results of nutritional quality of the food samples are shown in Table 4. The total digestibility (TD), biological value (BV), protein retention efficiency (PRE), nitrogen retention (NR) and net protein utilization (NPU) are given as 97.76–98.00, 9.83–29.32%, 4.76–5.57, 7.31–8.03, 9.80–35.67, respectively. However, the food efficiency ratio (FER), protein efficiency ratio (PER), net protein retention (NPR) are 0.07–0.08, 0.58–1.30, 0.33–0.34, respectively. All the experimental formulated diets had significantly (p < 0.05) nutritional (protein) qualities similar to the commercial control (cerelac) with the exception of BV and NPU. However, the BV and NPU of the experimental diets are significantly (p < 0.05) higher than those of basal diet (Table 4). The differences in the nutritional (protein) qualities of dried germinated QPM (DGQPM) and commercial cerelac might be due to the differences in their protein contents (~ 26 and ~ 23%), respectively. Interestingly, the BV of the current blends (10–29%) did not fall within the ranges (70–100%) reported for protein materials to be used as adequate and good nutritional quality sources for complementary food formulation (Ijarotimi and Keshinro 2013). This could be attributed to the complex metabolic process of the almond seed-enriched blends during which lipids, carbohydrates and storage protein within the seeds were broken down and prevented the bioavailability of the required protein (Jachmanian et al. 1995). Moreover, the
current BV in this study is comparatively low to the 41–60 (Jiarotimi and Keshinro 2012a, 56.16–89.92 (Abiose et al. 2015) and 88.31–100% (Akinsola et al. 2017) previously reported for different formulated complementary foods from germinated popcorn-bambara groundnut-African locust bean and quality protein maize-common maize, respectively.

### Hematological characteristics of rats fed with the formulated flour blends

The result of hematological characteristics of rats fed with the blends of quality protein maize (QPM) and almond seed flours is shown in Table 5. The packed cell volume (PCV), white blood cells (WBC) and platelets (RBC) of the rats fed with the flour blends were 31.00–49.00, 30.2–424 × 10³, 17.14–228 × 10³, respectively. Meanwhile, the other blood components such as neutrophil, eosinophil, basophil, monophil and lymphocytes were obtained as 43.40–53.4, 0.60–1.6, 0.01–0.8, 0.2–0.6 and 46.00–53.20, respectively. The different processing methods employed in the production of these formulated blends significantly \(^{(p < 0.05)}\) improved the haematological data of the experimental rats. For instance, the data (Table 5) showed that rats fed with the fermented high QPM (FHQPM) and dried germinated QPM (DGQPM) have significantly \(^{(p < 0.05)}\) higher PCV (49 and 47.40%) and RBC (228 and 280 × 10³) but lower WBC (42 and 30.2 × 10²) than the PCV, RBC and WBC of those fed with cerelac (47.20%; 280 × 10³; 51 × 10⁵), respectively. Notably, the rats fed with FHQPM, which had better PCV and RBC than those fed with DGQPM, showed that the fermentation process improved the blood formation ability of the blends to prevent the pathogenesis of anaemia. This observation is contrary to the protein content improvement observed with germination processing method of producing the blend (Table 1). Meanwhile, a previous study had reportedly linked the association of low PCV, RBC and serum protein with protein deficiency (Aleotor and Egbefon 1992). Thus, the high PCV and RBC but low WBC of the rats fed on the current blends relative to cerelac showed that the blends possessed the greater potentials of blood,

### Table 5 Hematological characteristics of rats fed with the blends of maize and almond seed flour

| Hematological characteristics | Samples | DQPM | DGQPM | FYM | FHQPM | CERC | BASAL |
|------------------------------|---------|------|-------|-----|-------|------|-------|
| PCV (%)                      |         | 36.40 ± 5.07^{d} | 47.40 ± 11.32^{ab} | 31.00 ± 12.14^{c} | 49.00 ± 7.61^{a} | 47.20 ± 4.15^{ab} |
| WBC (×10³/mm³)               |         | 3.20 ± 0.20^{b} | 3.00 ± 0.90^{b} | 3.00 ± 0.50^{b} | 4.20 ± 1.10^{ab} | 5.10 ± 1.20^{a} |
| PLATELETS (×10³/mm³)         |         | 190.00± | 280.00± | 171.00± | 228.00 ± 36.90^{a} | 208.00 ± 34.43^{c} |
| NEUT (%)                     |         | 43.40 ± 3.23^{c} | 51.80 ± 3.64^{a} | 53.40 ± 3.58^{a} | 47.00 ± 2.18^{b} | 49.00 ± 2.00^{ab} |
| EOS (%)                      |         | 1.60 ± 1.14^{a} | 1.60 ± 1.14^{a} | 0.60 ± 0.01^{a} | 1.20 ± 0.04^{b} | 1.00 ± 0.01^{c} |
| BASO (%)                     |         | 0.80 ± 0.05^{a} | NIL             | 0.40 ± 0.05^{b} | 0.20 ± 0.01^{c} | NIL             |
| MONO (%)                     |         | 0.60 ± 0.05^{a} | 0.20 ± 0.05^{b} | 0.20 ± 0.02^{b} | 0.60 ± 0.04^{a} | 0.20 ± 0.05^{b} |
| LYMPH (%)                    |         | 53.20 ± 7.30^{a} | 46.00 ± 7.50^{b} | 47.40 ± 4.72^{a} | 53.60 ± 4.04^{a} | 48.80 ± 3.27^{b} |

DQPM: Dried Quality Protein Maize, DGQPM: Dried Germinated Quality Protein Maize, FYM: Fermented Yellow Maize, FHQPM: Fermented High Quality Protein Maize, CERC: Cerelac, PVC: Packed Cell Volume, WBC: White Blood Cell, NEUT: Neutrophil, EOS: Eosinophil, BASO: Basophil, MONO: Monocytes, LYMPH: Lymphocytes. Values with different superscript on the same row are significant (p < 0.05).
antibody and cell-mediated immunity formation (Ijarotimi et al. 2015). Furthermore, previously reported findings (Ijarotimi and Keshinro 2012b; Abiose et al. 2015) had supported the above observation on flour blends from plant origins to contribute to the good haematological indices (high PCV and RBC with low WBC) of the fed subjects. However, there exist no significant differences ($p > 0.05$) in the other indices (neutrophil, eosinophil, basophil, monophil and lymphocytes) of the rats fed with the formulated diets relative to cerelac. This implied that the formulated diet did not contain any possible antinutritional factors (such as phytate, oxalate, etc.) above the health-hazardous level that could adversely affect the haematological and immune status of the animals (Ijarotimi et al. 2015).

**Effect of the formulated complementary diets on the quality and development of the internal organs of the experimental animals**

The protein (nutritional quality) of the internal organs of the animals is a determinant of the physiological needs of the rat, which denoted their biochemical (haematological) indices and resultant growth and development patterns of the animals after feeding with the formulated diets. The result of the effect of the complementary diet on the quality of the internal organs of the experimented animals is presented in Fig. 1. The result showed that the protein contents of the internal organs (liver, kidney and tissue) of the experimental animals (45.19–49.49, 45.85–49.21 and 46.56–51.88 mgN/g) after feeding with formulated blends were significantly ($p < 0.05$) lower than those (55.04, 55.91 and 53.72 mgN/g) of rats fed with the cereals (commercial control) respectively. Notably, the low BV (10–37%) and NPU (10–36) of the complimentary diets (Table 4) when compared to BV (70%) and NPU (68) of cerelac, even with similar digestibility (~98%) might have contributed to the low protein contents of the internal organs of the formulated diets-fed rats.

However, the growth and development of the internal organs (kidney, liver and heart), which could be either larger (overweight) or smaller (underweight) than the normal (control) sizes, depicted the growth patterns and health status of the animals. Therefore, the result of the effect of the complementary diet on the development (weight) of the internal organs of the experimental animals is presented in Fig. 2. The weights of the internal organs (liver, kidney and tissue) of the animals fed with experimental diets (1.36–2.7, ~0.54 and 0.25 g) were similar when compared to those fed on cerelac (1.98, 0.58 and 0.25 g), respectively. The present study is contrary to the past finding (Ijarotimi and Keshinro 2012b) that reported less weights of the internal organs of the animals fed with fermented popcorn-African locust bean-Bambara groundnut flour blends than compared to cerelac. Although, the rats fed with dried germinated QPM (DGQPM) sample have better protein contents and weights of internal organs when compared with those fed on fermented high QPM (FHQPM) sample but the current data generally revealed that the formulated diets did not result in abnormal development of the vital organs of the rats. Similar observations of no-toxic and high-safe consumption effects were previously reported on the rats fed with soybean-cowpea tempe-maize-based (Osundahunsi and Aworh 2003),
cassava-based (Onabanjo et al. 2008) and unripe plantain-soybean cake-rice bran-based (Odebode et al. 2017) complementary diets.

Sensory attributes of the formulated complementary diets

The sensory attributes of the complimentary diets presented in Table 6 showed that the panelists generally accepted (7.00–8.01) all the samples that were produced from the almond-maize flour blends except for the aroma of the products, which were neither liked nor disliked (5.00–5.88) by the panelists. Notably, the panelists did not report any significant difference \((p > 0.05)\) between the colour of the control (cerelac) and germinated (DGQPM) samples. The reason could have been that most of the minor components such as phenolic compounds might have been leached out during the germination process of producing DGQPM, thereby helping to improve its colour. Although, the control (cerelac) sample had a significantly \((p < 0.05)\) better texture than all the formulated diets but there was no observable difference in the consistency of the control, DQPM and DGQPM samples. Past studies on the maize-based snacks (Arise et al. 2018) and custard (Akinwale et al. 2017) were in agreement with the current work, which revealed no negative effect on the sensory (organoleptic) attributes of the complimentary diets after supplementation with high protein legume/oil crops, such as Bambara groundnut and soybean, respectively.

Conclusion

The study revealed that the complementary food formulated from almond seed, yellow maize and high quality protein maize compared favorably with cerelac to meet the nutritional needs of children above six months of age. The dual germination and fermentation methods employed during the processing stages thus, improved the nutritional qualities (protein contents, digestibility and amino acid composition) and haematological indices of the formulated complementary foods, respectively.

Table 6 Sensory attributes of the complementary diets

| Sensory attributes | DIET SAMPLES | DQPM | DGQPM | FYM | FHQPM | CERELAC |
|--------------------|--------------|------|-------|-----|-------|---------|
| Colour             |              | 6.39 ± 0.50<sup>b</sup> | 6.77 ± 0.80<sup>a</sup> | 6.17 ± 0.38<sup>c</sup> | 6.28 ± 0.46<sup>c</sup> | 6.78 ± 0.80<sup>a</sup> |
| Taste              |              | 6.61 ± 0.69<sup>b</sup> | 6.28 ± 0.46<sup>d</sup> | 6.33 ± 0.48<sup>b</sup> | 6.50 ± 0.51<sup>b</sup> | 7.61 ± 0.70<sup>a</sup> |
| Aroma              |              | 5.88 ± 0.83<sup>a</sup> | 5.11 ± 0.32<sup>d</sup> | 5.00 ± 0.02<sup>d</sup> | 5.33 ± 0.48<sup>b</sup> | 5.89 ± 0.85<sup>a</sup> |
| Consistency        |              | 6.39 ± 0.69<sup>a</sup> | 6.33 ± 0.48<sup>a</sup> | 6.11 ± 0.32<sup>d</sup> | 6.00 ± 0.03<sup>b</sup> | 6.40 ± 0.70<sup>a</sup> |
| Texture            |              | 6.72 ± 0.89<sup>a</sup> | 6.22 ± 0.42<sup>c</sup> | 6.11 ± 0.32<sup>d</sup> | 6.00 ± 0.02<sup>d</sup> | 7.73 ± 0.90<sup>a</sup> |
| Overall acceptability |            | 7.00 ± 0.02<sup>b</sup> | 7.00 ± 0.02<sup>b</sup> | 7.00 ± 0.03<sup>b</sup> | 7.00 ± 0.01<sup>b</sup> | 8.01 ± 0.02<sup>a</sup> |

DQPM Dried Quality Protein Maize, DGQPM Dried Germinated Quality Protein Maize, FYM Fermented Yellow Maize, FHQPM Fermented High Quality Protein Maize. Mean values with different superscript on the same row are significant \((p < 0.05)\)
The low swelling index, water absorption capacity, loose and bulk density values in this work indicated that higher amount of the flour particles could bond together and increase the beneficial effect of energy contents of these diets. Thus, it could be inferred that the functional properties of these diets would lead to production of appropriate complementary diets with adequate texture, dietary bulk and caloric density. The significant similar quality and weights of the internal organs of the experimental animals after feeding with formulated blends and cereals revealed their safe consumption with no possible adverse health effects, thus adding value to the locally produced almond seeds. Moreover, the improved nutritional qualities of the almond-quality protein maize-yellow maize flour blends therefore, encouraged its utilization in the global preparation of future complimentary foods for young children.

Abbreviations
BASO: Basophil; BV: Biological value; CERC: Cereals; DGQPM: Dried germinated quality maize protein meal; DJQPM: Dried quality protein maize; EDTA: Ethylene diamine tetrachloroacetic acid; EOS: Eosinophil; FER: Food efficiency ratio; FHPQPM: Fermented high quality protein maize; FYM: Fermented yellow maize; LYMHP: Lymphocytes; MONO: Monocytes; NEUT: Neutrophil; NPR: Net protein retention; NPU: Net protein utilization; NR: Nitrogen Retention; PER: Protein efficiency ratio; PRE: Protein retention efficiency; PVC: Packed cell volume; QPM: Quality protein maize; RDA: Recommended daily allowance; TARAA: Total aromatic amino acid; TDA: True digestibility; TEAA: Total essential amino acid; TSM: Technicon Sequential Multi-Sample; WBC: White blood cell

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Authors’ contributions
Adesanmi, R.A. collected and analyzed the data as well as prepared the draft of the manuscript. Malomo, S.A. thoroughly read and edited the manuscript as well served as the corresponding author of the manuscript. Fagbemi, T.N. thoroughly read and edited the manuscript. Adesanmi, R.A. collected and analyzed the data as well as prepared the draft of the manuscript. Malomo, S.A. thoroughly read and edited the manuscript. Fagbemi, T.N. also served as the corresponding author of the manuscript. Fagbemi, T.N. thank the laboratory staff of the department of food science and technology, Federal University of Technology, Akure, Nigeria.

Competing interests
The authors declare that they have no competing interests.

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