

Nutritional Composition of Value-added Fish Products from Selected Fish Species in Kenya

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Abstract

Fish is an important seafood that provides quality nutrients to human beings which is vital for health and development. This is especially critical in the diets of children under two years old and reproductive women (pregnant and lactating). Selected fish species were sampled from both Lake Victoria and fish farms, and their weight was determined using a sensitive weighing balance and fish-based products (powder and gelatin) developed to food-grade standards. The proximate composition of minerals, amino acid, and fatty acid profile were assessed in the laboratory, and data were analyzed using R statistic tool version 4.2.2 to understand the differences between the variables at a significance level of $p < 0.05$. Results indicated that crude fat ranged from 3-23%, crude protein was highest in Nile perch gelatin at 81.22 ± 1.43 while *omena* and *haplochromine spp.* had the highest concentration of calcium at 6347.1 ± 428.5 and 4522.0 ± 233.3 mg/Kg respectively. The highest concentration of essential amino acids was found in *omena* and Nile perch powder. Among the fatty acids; oleic and palmitic were the dominant fatty acid in the fish products at $25.12 \pm 0.56\%$ - $45.12 \pm 0.65\%$ and $20.52 \pm 0.69\%$ - $44.23 \pm 0.74\%$ respectively. The moisture content of the final products was between $4.64 \pm 0.08\%$ - $8.82 \pm 0.35\%$, which was within limits ensuring little microbial activity thus enhancing the shelf-life of the final products. The developed products had nutrients that are beneficial to human health at desirable concentrations and are recommended for inclusion in the diet of vulnerable groups for a prosperous community free from hunger and malnutrition.

Keywords: proximate composition, minerals, amino acid, fatty acids profile

1. Introduction

Fish is vital for proper human nutrition providing approximately 17% of the global animal protein intake (FAO, 2020). Fish is rich in important micronutrients, minerals, vitamins, amino acids, and essential fatty acids which are often lacking in the diets of poor and vulnerable populations (Thilsted et al., 2016) especially children below 2 years of age, and reproductive women (pregnant and lactating). In addition, this vulnerable population rarely consume fish mainly due to affordability and availability of convenient products. Therefore, the development of nutrient-rich products from fisheries and aquaculture contributes to food and nutrition security for many households thereby meeting the nutrient requirements for optimal growth and development (Gibson et al., 2020). This is mostly important for pregnant and lactating mothers, and infants as the first 1000 days are the most critical days of their life (Longley et al., 2014; WorldFish, 2017). Poor nutrition emanating from chronic malnutrition and micronutrient deficiency during these days can result in permanent and irreversible effects (Victoria et al., 2008) with a cost on the health and performance of affected individuals (Grantham-McGregor et al., 2007).

Undernutrition which is evidenced in stunting, wasting and underweight can result to disability or even death. Forty-five percent of mortality in children under five is attributable to under-nutrition; nutritional deficiencies are

responsible for 50% of years lived with disability in children aged four and below (Golden et al., 2016). This condition is evident in many regions in the globe and has continued to rise in Africa from 50 million people in 2000 to 59 million people in 2018 (UNICEF, 2019). In Kenya, 18% of children below 5 years are stunted, 5% are wasted while 10% are underweight (KNBS and ICF, 2023). It is therefore important to feed this vulnerable population with the right diet because the nutrition of the mother and the child significantly affects the child's future well-being and development which further affects educational and economic achievements. (SSentongo, 2021).

Fish are an important source of essential amino acids, long-chain polyunsaturated fatty acids, micronutrients such as vitamins A, B12, and D, and minerals essential for the cognitive development of children and for adult health (Golden et al., 2016; FAO, 2018). In addition, fish is recognized for its importance in enhancing the bioavailability of nonheme iron and zinc from plant-based calorie-dense foods such as rice consumed in the same meal (Beveridge et al., 2013). However, many Kenyans especially pregnant, lactating mothers and children below 2 years old consume lower fish servings than recommended levels by the World Health Organization (WHO) depriving them of vital benefits important for proper growth and development (WorldFish, 2017). In addition, during pregnancy and lactation, women have a higher demand for essential nutrients present in fish products. Poor fetal growth and development resulting from micronutrient inadequacy make children susceptible to opportunistic infections.

Despite the rich nutritional value of fish, intake by this vulnerable age group remains low, especially among the poor due to challenges with the availability and accessibility of convenient products, especially for children. This coupled with an increase in population (KNBS, 2020), has caused a general reduction in the frequency of fish consumption within households further reducing the national fish intake levels when compared to the rest of the World (FAO, 2020). Fish consumed in Kenya is predominantly provided by capture fisheries sourced from rivers, large inland lakes, coastal systems, and aquaculture (Munguti et al., 2021). The major fish species cultured in Kenya are Nile tilapia (*Oreochromis niloticus*) and African catfish (*Clarias gariepinus*) while the main source of capture fisheries is Lake Victoria with three major species being Nile perch (*Lates niloticus*), Nile tilapia and silver cyprinid/omena (*Rastrineobola argentea*). Other species such as Lung fish (*Protopterus aethiopicus*), African catfish, and *Haplochromines spp.* form an important component of the local diet among the riparian communities.

This study sought to develop and assess the nutritional composition of value-added fish products, namely fish powder and fish gelatin for children below 1000 days old and reproductive women throughout the year for food and nutritional security.

2. Materials and Methods

2.1 Sample Collection and Processing of the Value-added Fish Products

A total of six different fish species were collected for the preparation of value-added fish-based products, two species (silver cyprinid/omena and haplochromine spp) constituted the small fish food that is eaten whole. The big fish collected were Nile tilapia (500-800g), Nile perch (2000-3500g), African catfish (1000-3000g), and Lung fish (1500-3000g). Omena, haplochromine, Nile perch and Lung fish were collected from Lake Victoria, Dunga beach in Kisumu County while Nile tilapia and African catfish were collected from a fish farm in Kakamega County.

Gelatin was extracted through heat hydrolysis from Nile perch and Nile tilapia skin. The skin was cut into small pieces to increase the surface area-to-volume ratio for maximum extraction and to improve the efficiency of the extraction process. The resulting sap was then concentrated at 60⁰ C before molding into sheets and milling into gelatin powder.

To prepare fish powder from the big fish, the fish were descaled, gutted, washed to food grade standards, and cut into desirable small pieces. Deboning was done to separate the fillet from the adhering bone and partial boiling was done for five minutes at 60⁰ C. The separated fillet was used to make fish powder. The moisture content was reduced to below 10% and milling was done to produce the final product.

2.2 Laboratory Analysis

The samples were packed into labeled sterilized air-tight containers and stored in the freezer at -20⁰ C for laboratory analysis. All samples were analyzed in triplicate.

2.3 Proximate Analysis

The moisture content was determined by evaluating the loss of weight after drying the sample in a hot air oven at 105⁰ C for 3 hours until the samples achieved a constant weight as per the (AOAC, 2005) method. Crude lipid was determined by extracting fat using the Soxhlet method (AOAC, 1990). Ash content was determined in moisture-free dry samples in a muffle furnace at 550⁰ C for 20 hours until all organic components of the sample were

incinerated completely (AOAC, 1990). The crude protein content of samples was determined by multiplying the nitrogen content obtained by Kjeldahl's method by the conversion factor of 6.25 (AOAC, 1990). The crude fiber content of the sample was determined according to (AOAC, 1980); whereby 2 grams fat-free sample was digested with 1.25% (0.26 N) H₂SO₄ and 1.25% (0.23 N) NaOH and followed by drying in an oven at 105° C and then for 3 hours in a muffle furnace at 550° C. The energy value was determined by multiplying the crude protein, carbohydrate, and crude lipid content of the sample as per the method described by Jabeen and Chaudhry (2011). Total carbohydrates or nitrogen-free extract (NFE) of the sample were calculated by subtracting the sum of percentages of moisture, crude protein, ash, crude lipid, and crude fiber in the sample from 100% (Castell and Tiews, 1980).

2.4 Mineral Analysis

Mineral content was determined using 5 grams of the sample according to AOAC (2005). Samples were dissolved and digested in 15 ml HNO₃ which was topped up to 50 ml with distilled water after digestion. The aliquot was filtered using Whatman paper (0.45µm thickness) after which the samples were injected and determined using the atomic absorption spectroscopy method (Model AA-7000, Shimadzu, Corp., Kyoto, Japan) (Gokoglu et al., 2004). Potassium and phosphorus contents were analyzed by using the flame photometry method (Junsomboon and Jakmunee, 2011). Iodine was analyzed using the titrimetric/iodometry method.

2.5 Amino Acid Profile

The profiling of amino acids was conducted using the high-performance thin-layer chromatographic (HPTLC) method (Nilima and Kunda, 2014). Samples were hydrolyzed in an acidic solution of HCl at 110° C for 24 hours. The samples were then evaporated in a water bath to remove the trace acid content completely. The individual amino acids were quantified and identified based on the retention time and peak area of the amino acid's standards in HPTLC (Vijayan et al., 2016).

2.6 Fatty Acid Determination

The fatty acid profile was determined by gas chromatography. The extraction of lipids was done by modification of the Bligh and Dyer method (1959). 2 ml of water and 7.5 ml of methanol-chloroform (2:1 v/v) were added and the mixture was shaken overnight at room temperature. The supernatant was separated from the residue. The fatty acids were then obtained and converted into fatty acid methyl esters (FAMES) by methylation using HCl and methanol mixture. The composition and quantification of individual fatty acids were then determined by comparison of the retention times and peak areas with the respective peak areas of the relevant fatty acids' standards (Mohanty et al., 2019).

3. Statistical Analysis

Descriptive statistics were performed and results were presented as mean ± standard deviation (SD). Statistical analyses were conducted by using the r statistic tool version 4.2.2 to understand the differences between the variables at significance level of p < 0.05.

4. Results and Discussion

4.1 Proximate Composition

The moisture content of the prepared fish powder and fish gelatin was below 10% for all the products. This was important in reducing the microbial activities since most microbial activities are inhibited at moisture content below this level (Gram and Dalgaard, 2002) thus increasing the shelf-life of the final product (Dalgaard, 2000). The low moisture content observed in the final product was due to the processing technology which employed programmed drying of the samples that was preceded by milling to make the fish powder. Basiouny (2014) recorded a moisture content of between 9.2-11.9% in Nile tilapia when drying using the oven method (AOAC, 2005). Oduor-Odote et al. (2010) observed slightly higher values of final moisture in *omena* and *kimarawali* (*Stolephorus delicatulus*) at 12.1 and 14.0% respectively.

The crude fat content varied from 3.0 - 23.0%. The amount of fat in fish may be below 1% and may exceed 30% depending on the species and biological status (Cakli et al., 2007). Ash content varied from 1.7-16.7% with the highest value recorded on small fish food (silver cyprinid/*omena* and *haplochromine spp*). The high variation observed was due to the contribution of bones and scales since the small fish whole body was used in the preparation of the final product. Bille and Shemkai (2006) recorded ash content of 15.7% and 17.3% for spiced-smoked *omena* and sun-dried *omena* respectively. Bogard et al. (2015) reported 1.8% ash in Nile tilapia. Biswas et al. (2018) observed similar results at 1.83% ash in Nile tilapia collected from Dekar haor in Bangladesh. Likewise, Jim et al. (2017) found an ash content of 1.76 - 3.30% for Nile tilapia collected from different lakes in

Zimbabwe.

Protein was highest in Nile perch gelatin at 81.22 ± 1.43 while Lung fish and Nile perch values were 77.43 ± 1.21 and 75.53 ± 3.74 respectively. The lowest protein content was recorded in *omena* and *Haplochromines* at 55.1% and 56.2% respectively. Kabahenda et al. (2011) when assessing the protein levels in low-value fish, found a crude protein in the range of 47.9 - 58.8% in *omena*. In other studies, fillets from meager fish (*Argyrosomus regius*), the protein content was 75% of dry matter, the fat content was 19%, and the ash content was 5%. Corresponding values for gilthead sea bream (*Sparus aurata*) fillet were 57% protein, 40% fat, and 4% ash (Yiannopoulos et al., 1999). The value-added products (Nile perch skin and Nile tilapia skin) were also processed with the skin on hence may have contributed further to the protein. A study by Sinanoglou et al. (2014) on the by-products of meager fishes showed that the skin had the highest protein concentration. This explains the contribution of the skin to the protein of the final product.

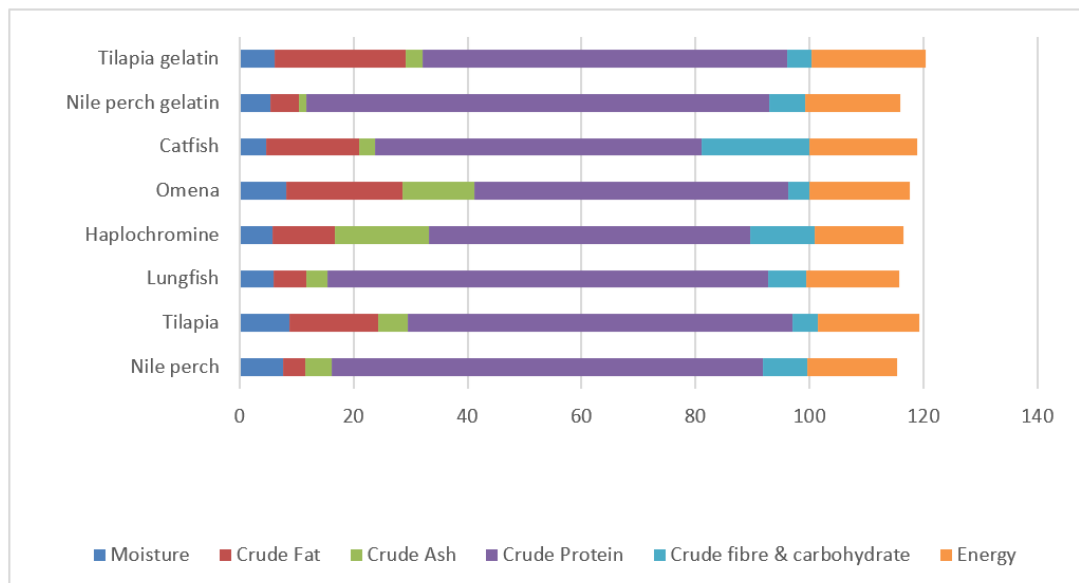


Figure 1. Bar graph showing percentage proximate analysis results of the value-added products from different fish species from Lake Victoria Basin

The high ash content and protein observed in this study could be attributed to the fact that the small fish were processed whole hence the bones and other hard tissues contributed to the high ash content (Usydus et al., 2011). Aquatic organisms take inorganic materials from food and water and store them in skeletal tissue and other membranes (Cakli et al., 2007). Generally, the proximate composition was significantly varied among the studied value-added fish products. This may be due to the absorption capability and conversion potentials of essential nutrients from their diets or their feeding niche (Adewoye and Omotosho, 1997).

4.2 Mineral Composition

Table 1 below shows the mineral composition of the fish powder and gelatin from Nile perch and Nile tilapia skin. All the minerals showed significant differences. *Omena* and *haplochromine spp.* had the highest concentrations of calcium (6347.1 ± 428.5 and 4522.0 ± 222.3 mg/Kg) respectively. The two species comprise the small fish species in Kenya and contribute more micronutrients since they are consumed whole (Thilsted et al., 2016; Roos et al., 2007). Small fish consumed whole are rich in bioavailable micronutrients such as zinc, iron, and calcium (Aakre et al., 2020) and are mostly nutrient dense. This explains why they have more nutrients than fillets. One study found that calcium absorption from the consumption of soft-boned small fish was comparable to that of skimmed milk (Hansen et al., 1998). Potassium levels in the six fish species ranged from 5030.8-9364.7 mg/Kg while phosphorus levels ranged from 1453.8-3163.7 mg/Kg. Jim et al. (2017) found potassium levels in Nile tilapia from three different lakes in Uganda ranging from 3193.3-4293.3 mg/Kg while phosphorus levels were ranging from 1453.8-3163.7 mg/Kg. Another study by Islam et al. (2021) recorded much lower levels of potassium in Nile tilapia in the range of 1096.5-1323.7 mg/Kg.

Trace elements (micro-nutrients) also varied significantly at $p < 0.05$. Zinc levels were highest in *omena* and

haplochromines spp at 67.5 and 69.6 mg/Kg respectively. Kabahenda et al. (2011) assessed micronutrient levels in *omena* from the Lake Victoria region and found zinc in the range of 41-103 mg/Kg on a dry basis. Clarke et al. (2022) found much higher levels of zinc at 170 mg/Kg from *omena* collected from three East African Lakes. The highest manganese content was observed in *omena* and *haplochromine spp*. at 11.5 and 26.5 mg/Kg respectively. Manganese levels in Nile perch and Nile tilapia was 2.01 and 2.61 mg/Kg respectively while it was not detected in catfish and Nile perch gelatin. Islam et al. (2021) reported that manganese content ranged from 1.02 mg/kg to 4.11 mg/kg in Nile tilapia.

Clarke et al. (2022) concluded that increasing the dietary intake of zinc, iron and calcium is important in curbing micronutrient deficiencies. Worth noting is that it is difficult to meet the Daily Recommended Intake (DRI) of these nutrients from plant-based food sources (Gibson et al., 2020) and therefore animal based protein sources such fish are key for women and children nutrition and wellbeing. Fish obtain minerals from their diet, feeds and also from weathered rocks and soils, and also from minerals present as an electrolyte (Tagliabue et al., 2017). Thus mineral absorption from the environment depends on concentration and species differences which in turn affects homeostasis (Lall, 2021). These variations in factors and conditions may explain the difference in the concentrations of these elements as observed in this study.

Table 1. The table shows the mineral concentrations of fish powder and gelatin processed from different fish species. ND indicates elements not detected

Essential minerals (macro-nutrients expressed in mg/Kg)				
	Calcium	Potassium	Phosphorus	
Nile perch	1842.8±227.4	9093.93±775.9	2584.8±226.3	
Nile tilapia	2382.4±151.9	7336.5±578.5	1668.4±77.4	
Lungfish	2236.2±278.5	7966.2±498.1	1757.9±44.6	
<i>Haplochromines</i>	4522.0±222.3	9364.7±428.3	2916.8±60.1	
<i>Omena</i>	6347.1±428.5	7181.3±557.9	3163.7±42.6	
Catfish	2447.4±37.7	8361.5±377.8	2157.8±124.1	
Nile perch gelatin	3841.5±24.18	5030.8±297.2	1453.8±131.4	
Nile tilapia gelatin	3362.1±16.4	6293.8±384.6	1789.8±104.0	
Trace elements (micro-nutrients expressed in mg/Kg)				
	Zinc	Iodine	Copper	Manganese
Nile perch	32.2±1.3	1.34±0.02	3.29±0.09	2.01±0.18
Nile tilapia	28.8±2.5	0.98±0.01	4.39±0.30	2.61±0.11
Lungfish	49.43±2.9	0.64±0.01	3.05±0.27	3.16±0.28
<i>Haplochromines</i>	67.5±2.6	0.70±0.02	7.32±0.20	11.48±0.16
<i>Omena</i>	69.6±3.3	0.28±0.01	5.15±0.49	26.54±1.65
Catfish	14.7±0.4	0.34±0.01	3.42±0.13	ND
Nile perch gelatin	7.1±0.5	0.22±0.02	7.45±0.45	ND
Nile tilapia gelatin	10.3±0.7	0.39±0.02	3.12±0.17	4.63±0.31

4.3 Amino Acid Profile

Fish are important sources of amino acids which are the building blocks for protein. Both essential and non essential amino acids were recorded from the samples. The essential amino acids included lysine, leucine, and isoleucine while the non essential amino acids included cysteine, tyrosine, and proline. Similar results were reported by Islam et al. (2021) who found essential amino acids in tilapia to be leucine and lysine while the non-essential were aspartic acid and glutamic acid. Similarly, Moses et al. (2018) reported that leucine, lysine, and aspartic acid to be the major amino acids in tilapia collected from different rivers in Nigeria. Additionally, the findings coincide with Adeyeye et al. (2018) who found aspartic acid and leucine to be the most common amino acids in tilapia. Kim and Lall (2000) investigated the amino acid contents of three kinds of marine flounder fish and found the most abundant amino acids to be aspartic acid, glutamic acid, glycine, leucine, and lysine. In another study, Mohanty et al. (2014) posited that small indigenous fish are rich in histidine while catfishes are rich in glutamic acid and glycine.

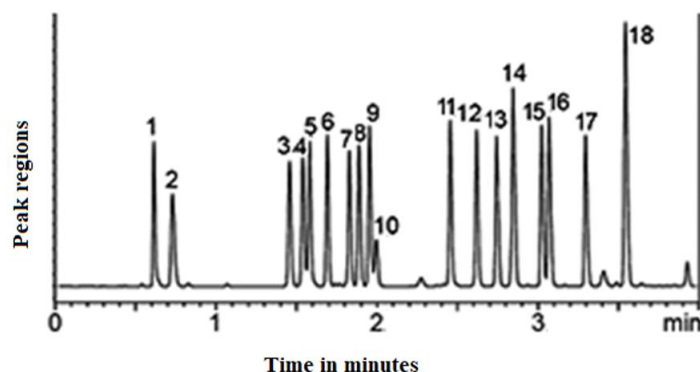


Figure 2. Shows the peak regions for different amino acid concentrations (1. Aspartate, 2. Glutamate, 3. Serine, 4. Glycine, 5. Histidine, 6. Arginine, 7. Threonine, 8. Alanine, 9. Proline, 10. Ammonia, 11. Tyrosine, 12. Valine, 13. Methionine, 14. Cysteine, 15. Isoleucine, 16. Leucine, 17. Phenylalanine, 18. Lysine)

The total essential amino acids (TEAAs) varied significantly at ($p < 0.05$) for all the products. However, there was no significant variation for specific essential amino acids throughout the products as shown in figure 3. Such differences could be attributed to differences in dietary needs of the respective amino acids, type of feed, and niche (Doğan and Ertan, 2017; Kuzu, 2005). It is also a reflection of differences in the protein composition of the product as amino acids are the protein building blocks (Abdullahi, 2002). The amino acid concentration determined in this study were within the standard amino acid values of FAO/WHO (1991) and values of FAO (2013). The products developed can therefore be depended upon for nutritional benefits.

Parameter1	Parameter2	r	95% CI	t(6)	p
Lysine	Threonine	-0.19	[-0.79, 0.60]	-0.47	> .999
Lysine	Histidine	-0.29	[-0.83, 0.52]	-0.74	> .999
Lysine	Isoleucine	0.21	[-0.58, 0.80]	0.53	> .999
Lysine	Leucine	-0.13	[-0.76, 0.63]	-0.31	> .999
Lysine	Methionine	-0.32	[-0.84, 0.49]	-0.84	> .999
Threonine	Histidine	-0.21	[-0.80, 0.58]	-0.52	> .999
Threonine	Isoleucine	-0.22	[-0.80, 0.57]	-0.56	> .999
Threonine	Leucine	-0.46	[-0.88, 0.36]	-1.26	> .999
Threonine	Methionine	-0.30	[-0.83, 0.51]	-0.77	> .999
Histidine	Isoleucine	-0.46	[-0.88, 0.37]	-1.25	> .999
Histidine	Leucine	-0.19	[-0.79, 0.59]	-0.47	> .999
Histidine	Methionine	-0.21	[-0.80, 0.58]	-0.53	> .999
Isoleucine	Leucine	0.63	[-0.13, 0.92]	2.00	> .999
Isoleucine	Methionine	-0.37	[-0.85, 0.45]	-0.99	> .999
Leucine	Methionine	-0.06	[-0.73, 0.67]	-0.15	> .999

p-value adjustment method: Holm (1979)
 observations: 8

Figure 3. Correlation of different essential amino acids showing the differences in their concentrations at $p < 0.05$

Amino acids are important biomolecules that serve as building blocks of proteins and are intermediates in various metabolic pathways (Mohanty et al., 2014). In addition, most of the animal sources are considered high quality protein with maximum indispensable amino acid composition for human needs and high digestibility. They are also precursors for the synthesis of various biologically important substances including nucleotides, peptide hormones, and neurotransmitters. Moreover, amino acids play an important role in regulating gene expression and cell signaling (Mohanty et al, 2014) and act as protein phosphorylation cascade (Wu, 2010), nutrient transport and metabolism in animal cells (Wang et al., 2013), and innate and cell-mediated immune responses.

Leucine, being the only dietary amino acid responsible for stimulating muscle protein synthesis (Etzel, 2004) also has important therapeutic functions in reducing stress conditions associated with burns, trauma, and sepsis (De

Bandt and Cynober, 2006). Arginine plays a critical role in cell division, wound healing, ammonia removal, immune function, and hormone release (Sharma et al., 2013). Methionine is key in improving wound healing, and is also used to treat liver disorders, depression and Parkinson's disease (Mischoulon and Fava, 2002). Amino acids such as glutamic acid, aspartic acid, alanine, and glycine are responsible for flavor and taste (Ruiz-Capillas and Moral, 2004) and therefore play a significant role in determining consumer taste preferences.

4.4 Fatty Acid Composition

The fatty acid profile was dominated by the Monounsaturated Fatty Acids (MUFA), Saturated Fatty acids (SFA), and Polyunsaturated Fatty Acids (PUFA) respectively. The most dominant SFA was palmitic acid. The highest concentration was recorded in *haplochromine spp.* while the lowest concentration was in lung fish. Previous studies also recorded palmitic acid as the highest among the SFA accounting for up to 65% (Huynh and Kitts, 2009; Usyduş et al., 2011; Zhang et al., 2020) in fish. Islam et al. (2021) documented SFA, MUFA and PUFA in Nile tilapia. Oleic acid was the highest MUFA in all the samples and the highest levels were recorded in Nile perch gelatin, African catfish powder, and Nile tilapia powder respectively. Mohamed and Al-Sabahi (2011) found palmitic, stearic, oleic and decosahexaenoic fatty acids present in Nile perch fillets. Li et al. (2011) reported a higher content of oleic acid but a lower content of palmitic acid in marine fish species from the East China Sea. Our findings also concur with research by Zhang et al. (2020) who found a similar trend in *Sillago sihama*, *Collichthys lucidus*, *Coilia mystus*, *Harpadon nehereus*, and *Johnius belangerii* while other species in the same study showed elevated SFAs, MUFA and PUFA respectively.

The products prepared in this study had the lowest SFAs and were healthy and potential in preventing malnutrition and cardiovascular diseases (Islam et al., 2021). Among the PUFA, linoleic acid, a metabolic precursor of arachidonic acid was the highest among all the samples followed by linolenic acid, a metabolic precursor of Eicosapentaenoic acid (EPA) and Docosahexaenoic acid (DHA). Saturated fatty acids showed a linear relationship with MUFA and an inverse correlation with PUFA. Figure 4 below shows the relationship between the fatty acids. This inverse relationship was also observed between MUFA and PUFA in the products developed. The high concentration of linoleic acid could be due to the fish diet and the retention of arachidonic acid in the fish tissue. Arachidonic is preferentially retained in the muscle's lipids while EPA is selectively used as a substrate for beta-oxidation hence resulting in low levels. (Suloma et al., 2008; Karapanagiotidis et al., 2010).

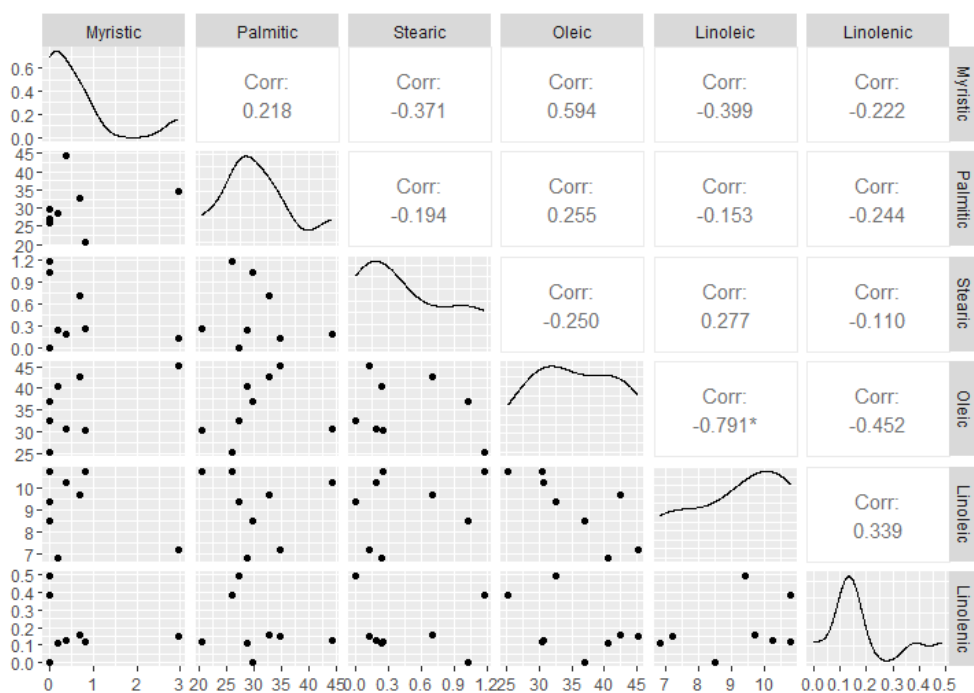


Figure 4. Correlation analysis of different fatty acids and their concentration in fish powder developed from common fish products

5. Conclusion

Value addition of fish through the preparation of fish-based products such as gelatin and fish powder is key in

minimizing post-harvest losses and increasing productivity. In addition, it ensures that fish products are available throughout the year and are easily accessible whenever needed. Considering fish central role in healthy diets, it is important to have nutrition sensitive interventions geared towards alleviating the prevalence of undernutrition such as stunting and wasting in children and micro nutrient deficiencies. Healthy fish diets are key in improving survival, health, physical growth, cognitive development and productivity in children and reproductive women. The fish based products developed were nutrient dense and therefore favourable for good health and development of the targeted vulnerable population. Most importantly, the nutrients were at recommended levels to meet the targeted population's requirements for optimal growth and development to create a healthy society free from hunger and malnutrition. The moisture content of the final product was low enough for minimum microbial activity thus ensuring a longer shelf life.

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Authors contributions

Study Conceptualisation: Cecilia Githukia, Dennis Otieno and Kevin Obiero. Sample preparation: Dennis Otieno, Evans Menach, Cecilia Githukia, Maureen Cheserek and Kevin Obiero. Data Collection: Domitila Kyule-Muendo, Dennis Otieno and Maureen Cheserek. Study design and revision: Jonathan Munguti, Evans Menach and Kevin Obiero. Drafting of the manuscript: Dennis Otieno and Cecilia Githukia. All authors read and approved the final manuscript. In addition, all authors contributed equally to the study.

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Competing interests

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

Informed consent

Obtained.

Ethics approval

The Publication Ethics Committee of the Canadian Center of Science and Education.

The journal's policies adhere to the Core Practices established by the Committee on Publication Ethics (COPE).

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Data availability statement

The data that support the findings of this study are available on request from the corresponding author. The data are not publicly available due to privacy or ethical restrictions.

Data sharing statement

No additional data are available.

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