

Review Article

Changes in Fish Quality During Canning Process and Storage Period of Canned Fish Products

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Abstract

Fish and fish products are subject to enzymatic, microbiological and chemical changes during fish processing. Canned fish products stable at ambient temperature, have long shelf life and in consequence are eminently suitable for world-wide distribution. Therefore this study aimed to conclude the alterations in proximate chemical composition, Quality characteristics, nutritional quality and microbiological safety of fish during canning process. Also following changes in these parameters during storage period of canned fish.

Keywords: Canning Process; Chemical Composition; Storage Period

Abbreviations: TVB-N: Total Volatile Basic Nitrogen; TMA-N: Total Trimethyleamin Acid Nitrogen; PV: Peroxide Value; NPU: Net Protein Utilization; BV: Biological Value; FAD: Food and Agriculture Organization; FDA: Food and Drug Administration.

Introduction

Canning is a well-established and traditional means of providing food which is stable at ambient temperatures, has long shelf life and in consequence is eminently suitable for world-wide distribution. Canned fish is therefore exported from countries all over the world into the consumer markets. The manufacturing of canned fish has provided, and continues to provide much-needed employment, individual incomes and the means for foreign currency exchange for developing countries, particularly in Southeast Asia, South America and the Indian Ocean [1]. Canned fishery products are formally canned in vegetable oils and now available in water packs. Preservation methods such as freezing and canning are technologies that used basically due to cost and non-availability of equipment and cold storage system [2]. The advantages of canned food are such as their preservation, safety and convenience. The major steps in canning process include cooking, cooling, packing with a covering oil or tomato sauce in sealed cans and sterilized to achieve commercial sterilization by the heat. Oxidation of lipids and formation of nitrogenous compounds can occur during the cooling step of fish cooking [3]. During storage temperature (above 35 °C) must be avoided to prevent the growth of thermophilic bacterial spores. Changes in the food quality during storage period result from a changes in physicochemical and microbiological properties that decrease their nutritional value, palatability, and safety.

Chemical Composition

Slabyj and Carpenter [4] stated that raw mussel meat contained 81.2% moisture, 3.29% protein, 0.81% lipid, 0.41% ash, and 0.75% carbohydrate, steamed mussel meat contained 74.6% moisture, 2.73% protein, 0.62% lipid, 0.24% ash, and 0.32% carbohydrate, and canned mussel meat contained 75.7% moisture, 2.90% protein, 0.72% lipid, 0.27% ash, and 0.20% carbohydrate. El-Sherif [5] stated that

the moisture contents of shrimp canned with vegetables and those canned with tomato sauce were lower by 12.6 and 10.8% respectively as compared with moisture content of raw shrimp. Also he found that protein content in shrimp and found that protein contents of raw shrimp, samples canned with vegetables and samples canned with tomato sauce were 76.90, 60.85 and 68.11 (on dry weight basis), respectively. These values indicated that heat treatments applied in canning process resulted in losses of protein which were accounted by 20.9 and 11.4 in samples canned with vegetables and tomato sauce respectively.

Chemical analysis of canned shrimp showed that crud fat; crud ash and carbohydrate content were higher than of raw shrimp [5]. The chemical composition of smoked mussels of (*Mytilus galloprovincialis, L.*) were 20.4 ± 0.45 , 4.9 ± 1.30 , 2.4± 0.02 and 69.4 ± 0.23 for Protein, lipid, ash and moisture respectively. After caning process these values changed to 12.7 ± 0.46, 8.1 ± 0.14, 3.1 ± 0.14 and 70.2 ± 0.62 for Protein, lipid, ash and moisture respectively [6]. Also he reported that chemical composition of mussels (Mytilus galloprovincialis, L.) were 17.3 ± 0.04, 4.0 ± 0.28, 1.8 ± 0.19 and 73.0 ± 0.07 , after caning process these values changed to 12.2 ± 0.47 , 10.6 ± 0.57 2.8 ± 0.01 71.1 ± 0.29 for Protein, lipid , ash and moisture, respectively. Shakila, et al. [7] have observed a reduction of 6% moisture content in cooked tuna. Also Garcia-Arias, et al. [8] have reported that moisture content decreased and fat content increased in cooked fish fillets. Selmi et al. [3] the lipid contents found in tuna (Thunnus thynnus) and sardine (Sardina pilchardus) contents dropped significantly (P<0.05) after the caning process. In cooked tuna, the moisture content decreased significantly (p<0.05)with an increase in the duration of heat process giving up to 16% reduction [9].

Canning process reduced the protein content for Orcynopsis unicolor, but increased in protein content for Euthynnus affinis. El-Dengawy, et al. [10] determined the chemical composition of sixteen samples of canned fish products imported (canned tuna, canned sardine, canned Mackerel) and observed that moisture percentages in all canned fish samples ranged between 52.41±0.035 to 78.53±0.142%. It could be observed that all canned fish samples had high values of water activity (0.990-0.999). Meanwhile, NaCl content ranged between 0.13±0.000 to 1.20±0.042 %. Czerner, et al. [11] determined that the chemical composition of fresh anchovy were 77.65±0.67, 16.24±0.82, 4.25±0.09 and 1.16 ± 0.06 for moisture content, protein content, lipid content and ash content respectively. In canned fish, chemical composition were 64.82 ± 0.06 , 26.74 ± 0.74 , $6.23 \pm$ 0.38 and 2.21 ± 0.01 respectively. Sajib [12] Studied the effect of canning process on the chemical composition of chela (Laubuka dadiburjori) and determined that moisture content, protein content, lipid content, ash content and carbohydrate

content of fresh fish were 76.56±1.62, 13.74±1.22, 4.25±0.85, 2.37±0.56 and 1.41±0.79 respectively. After canning process these values changed to 67.15±1.69, 16.68±0.88, 5.46±0.34, 8.15±0.83 and 1.35±0.07 for Moisture, Protein, Lipid, Ash t and Carbohydrate respectively. Fatma Arfat [13] found that the moisture, sodium chloride and protein contents of canned sardine and mackerel were decreased after 24 months storage at room temperature while, total lipids contents was increases. Ghaly [14] pointed out that chemical composition of canned shrimp was affected by storage at room temperature for 6 months and the kind of filling medium. Moreover, moisture, protein and carbohydrates were decreased while, fat and ash were increased in canned bosr and shrimp packed by dry method. On the other hand the moisture, fat and ash contents were increased [15]. Found that moisture contents of Tilapia fish fillets slightly decreased while crude protein contents was considerably reduced but crude fat and ash contents were progressively and gradually increased during storage of ambient temperature. Ibrahim [16] mentioned that moisture, protein and fat contents of canned sardine showed a gradual decreasing as a period of storage at room temperature extended up to 12 months while, ash and salt contents showed a noticed increases. The chemical composition of canned chela (Laubuka dadiburjori) fish were 76.56±1.62, 13.74±1.22, 4.25±0.85, 2.37±0.56 and 1.41±0.79 at zero time of storage period. After 30 day of storage period these values changed to 68.13±2.06, 15.15±0.95, 5.51±0.56, 10.43±0.91 and 1.36±0.09. At the end of storage 60 days these values were 68.88±1.89, 15.62±0.45, 5.96±0.48, 12.60±1.12 and 1.39±0.07 for moisture content, protein content, lipid content, ash content and carbohydrate content respectively [12].

Quality Characteristics

Total Volatile Basic Nitrogen (TVB-N) and Trimethyleamin (TMA-N)

TMAO can be degraded during the thermal processing to disagreeable molecules such as TMA and DMA [17, 5]. The cooking process had no significant effect on the TVB-N levels (P<0.05), while TMA-N levels in sardine increased to reach 6.32 mg/100g. [3]. El-Dengawy, et al. [10] determined the quality characteristics in Sixteen samples of canned fish products imported (canned tuna ,canned sardine, canned Mackerel) and observed that TVN values in canned fish samples ranged between 7.01±0.254 mg N/100g sample to18.04±0.593 mg N/100g sample. The TVB-N in fresh chela (Laubuka dadiburjori) fish recorded 7.10 mg N/100g, these value increased to 15.50 mg N/100 g after caning process [12]. Increasing the total volatile basic nitrogen was attributed to the degradation of nitrogenous compounds resulting in an increase in both trimethylamin nitrogen (TMAN) and ammonia nitrogen (NH₃ N) which were considered the principle fractions of the TVBN [13].

The total volatile nitrogen contents of canned shrimp and bosr packed by dry method and in 3% salt solution, slightly increased during storage at ambient temperature for 6 months [14]. Abd El-Ghafour [15] recorded a gradual increase in the TVBN content of canned Tilapia fish fillets products as the storage period was extended. At the end of 12 months storage, samples steamed and canned in oil or tomato sauce showed an increasing of TVBN determined by 15.58 and 17.05% respectively while, the increasing rates were 18.71 and 20.61% in samples fried and canned in the same filling media respectively. Fatma Arfat [13] observed a gradual increase in TMA contents of canned mackerel and sardine samples as the time of storage at ambient temperature was extended which was attributed to trimethylnitrogen oxide that natural present in fish tissue which chemically reduced by SH group existing in fish protein or by zinc present in C-Enamle used for varnishing the cans. Ibrahim [16] found that the highest increased observed in TMAN content of sardine during storage was found in samples canned with sauce comparing with those packed in oil or spices. The total volatile basic nitrogen of canned sardine packed in different media gradually increased as the period of storage at room temperature was progressed [16]. In the first period of storage (3 months), TVB-N and TMA-N values increased significantly to attain 23.85 and 18.42 mg/100 g, and 7.95 and 8.11 mg/100 g in sardine and tuna, respectively, and remained statistically constant (P<0.05) until the end of the storage period [3]. The TVB-N value of canned chela (Laubuka dadiburjori) fish was 15.50 mg N/100 g, at zero time of storage period. It increased up to 17.68 mg N/100 g after 30 days. After that it continued to increase to 21.95 mg N/100 g after 60 days of storage period [12].

Thiobarbituric Acid (TBA) and Peroxide Value (PV)

Primary (peroxide value; conjugated dienes) and secondary (TBA and carbonyl values) lipid oxidation detections did not afford accurate methods for testing quality differences in canned products. Chia et al reported that canning process of rainbow trout pollok and shrimp resulted in reduction of thiobarbituric acid by about 50% of its initial value in the raw materials. The evaporation of water and loss of juiciness during cooking might have also contributed to an increase in TBA-RS values after cooking [18-20]. Koizumi, et al. [21] have also reported that TBA-RS values increased during cooking of fish at 100°C for 30 min, but they were below 1 µg of malonaldehyde/g of fat. Thiobarbituric acid determined as malonaldhyde slightly increased from 0.40 mg /kg of raw shrimp to 0.44-0.48mg /kg of the canned samples. This could be attributed to slight oxidation of unsaturated fatty acids found in shrimp during heat treatment of canning Process [5]. El-Dengawy, et al. [10] determined the quality characteristics in sixteen samples of canned fish products imported (canned tuna, canned sardine, canned Mackerel)

and observed that acid value of canned fish samples had the highest value of AV being 20.39 ± 0.000 mg KOH/g oil in CM.

It is well-known that the quality of canned products has a very close relationship with their lipid content and composition. Canned fishery products are especially susceptible to flavour and other changes due to the high levels of polyunsaturated fatty acids [22]. Ghaly [14] observed a gradual increase in TBA value of canned Suez Shrimp and elongated boor samples as the period of storage at ambient temperature was prolonged. Ibrahim [16] who noticed that TBA values of canned sardine products were gradually decreased as storage period extended and samples canned with sauce showed the lowest TBA values. Abd El-Ghafour [15] reported that a progressive increment in the TBA value of canned Tilapia fish fillets during storage at ambient temperature for 12 months. Selmi, et al. [3] studied the effect of local canning process and storage time (up to 6 months) on tuna and sardine canned in olive oil and tomato sauce and found that the PV and TBA index increased significantly in tuna.

рН

pH values of raw shrimp, canned shrimp with vegetables and canned shrimp with tomato sauce were 6.27, 6.61 and 6.48 respectively. The higher PH values observed in canned samples may be due to the formation and accumulation of some dibasic amino acid and volatile basic nitrogenous compounds such as NH₃ as a result of breakdown and proteolysis of proteins during heat treatment [5]. The pH value of fresh chela chela (Laubuka dadiburjori) fish was 6.8. Drop off pH value after caning process to 5.9 in canned fish [12]. Czerner, et al. [11] studied the effect of canning process on physicochemical of anchovy (Engraulis anchoita) and determined that pH value of fresh fish was 6.07, after canning slightly increased to 6.12 Fatma Arfat [13] observed that pH values of canned sardine and mackerel products were increased during storage at room temperature due to protein degradation into basic products such as ammonia, amines and hydrogen sulphide.

Ghaly [14] showed that pH values of canned Suze Shrimp and bosr in dry pack or in 3% salt solution were slightly decreased during storage at room temperature for 6 months to 6.25 and 6.30 under the same condition. Abd El-Ghafour [15] during storage of canned Tilapia fish fillets at room temperature, the highest pH value was found in samples fried and canned in tomato sauce. Ibrahim [16] noticed a significant increase in pH value of canned sardine packed in oil, while samples packed in sauce showed a decreasing trend as the period of storage at room temperature was extended. Sajib [12]. Studied the effect of storage period on the quality of canned chela (*Laubuka dadiburjori*) and found that the PH value 5.9, 6 and 6.2 for 0, 30 and 60 day of storage period.

Nutritional Quality

Fatty Acid

Aubourg, et al. [18] reported a slightly higher value in certain fatty acids of cooked tuna. Also observed a significant decrease in C20:5 and C22:6 on cooking of albacore tuna. The cooking process (caning) had a significant effect (P<0.05) on SFA composition in tuna flesh lipids. The content of C14:0, C18:0 and C20:0 significantly changed; however, unsaturated fatty acids remained constant after processing [3]. The SFA, MUFA and PUFA contents of raw tuna were 15.5%, 18.3% and 57.9%, respectively. The major SFA were palmitic (C16: 0), stearic (C18:0) and behenic (C22:0) acids. After caning process, the SFA contents varied from 17.4 to 11.0% with an increase in the duration of cooking.

Slightly higher values were noticed in most of the SFA in cooked tuna compared to raw tuna except those cooked for 30 min. However, there were significant losses (p<0.05) in C14:0, C16:0, C18:0 and C22:0 fatty acids in tuna cooked for 30 min. The total MUFA content increased with cooking time from 18.8 to 24.5%. This was mainly due to the increase in C24:1 fatty acid. The PUFA content in cooked tuna was more or less same as that of their raw counterparts. There was no significant decrease (p>0.05) in C20:5 fatty acid, however C22:6 suffered 21% loss (p<0.05). Losses were significant (p<0.05) with respect to C16:3, C18:2 and C20:2 fatty acids [9]. Czerner, et al. [11] determined the FAs profile of anchovy (Engraulis anchoita) samples taken during the canning process and also of the covering liquid taken after thermal treatment and observed that Steam-cooking mainly affected the MUFAs and ω -6 fatty acids, whose contents were significantly reduced after this operation (p50.01). This decrease could be related to leaching loss, as shown the reduction of fat content after this step and also to lipid damage due to high temperatures. Selmi, et al. [3] found that a higher content of C18:1w9 and C18:2w6 fatty acids were found in the canned samples following 3 and 6 months of storage at ambient temperature. Canned sardine and tuna were characterized by their richness in linoleic (10.98-11%) and oleic (45.3-44.95%) acids after 3 and 6 months of storage, respectively.

Amino Acid

Protein denaturation by heat does not necessarily cause nutritional loss. However, denatured proteins become more reactive and can be easily damaged by interacting with other constituents. The total free amino acids content decreased during tuna canning, especially if over-processing was employed. Domah, et al. [23] stated that fresh, canned and cooked, mussels are rich in terms of methionine, lysine, cystine, threonine, tryptophan, phenylalanine, tyrosine and arginine when compared to those of beef or FAO reference protein. Also have found that frying or boiling increases the nutritional value of mussels, as demonstrated by our amino acid composition results. It was also reported that the nutritional value of fried canned mussels was high. The analysis of nitrogen balance of raw or canned tuna protein did not show significant differences between raw and canned tuna in digestibility (DC), biological value (BV) or net protein utilization (NPU). The loss in available lysine in the canned tuna appeared to have no influence on nutritional quality [24].

Heat treatment of canning process may adversely affects the quality of food protein through it influence in the protein contents of the individual amino acids. The general trend that observed was slight decreasing in some amino acids in the canned samples. The loss particularly more observed in histidine which decreased by 16.1 to 17.9%, lysine and arginine also showed a noticed decrease estimated by 4.3-7%, 4.4-8.4 % respectively. Sulphur containing amino acids that are methionine and cysteine showed a reduction accounted by 1.9-3.8% and 20-4.9% respectively [5]. Gulgun, et al. [6] studied the effect of caning process on the smoked mussels of (Mytilus galloprovincialis, L.) and found that through the smoking and canning processes the relative changes in the content of free amino acids of canned smoked mussels resulted in levels that were reasonably comparable to those of steamed mussels. It's estimated that water conditions where the mussels were cultivated and the additives used for pre-processing before smoking and canning might have been responsible. Glutamic acid content of mussel meat of (Mytilus galloprovincialis, L.), before and after canning, was rather high. No negative effects on glutamic acid content due to either smoking or canning were found [6].

Minerals and Vitamins

Some loss in minerals (Na, K, Mg, P, Cu, Fe, Ca) from the muscle into the dipping medium occurs in canned tuna [25]. Gall, et al. [26] reported that a high fat content in the flesh produced a low loss in minerals, indicating a kind of interaction between both types of constituents. An advantage of fish canning is that bones become soft textured and thus edible, providing an important calcium source [27]. Mineral content of the raw tuna were 3.65±0.7, 124.9 ± 53.2, 116.55±2.6, 1424.75±3.6, 1.55±0.2, 0.65±0.1, 3.2 ±50 and 978.0 ±1.0 for Ca, Mg, Na, K, Zn, Cu, Fe and P respectively, while in steamed tuna these values were3.75±0.4, 95.35±3.8, 114.25±3.5, 1082.15±4.8, 1.55±0.1, 0.65±0.1, 2.85±0.2 and 835.35±4.0 0 for Ca, Mg, Na, K, Zn, Cu, Fe and P respectively [24]. The heat-labile vitamins thiamine, riboflavin, niacin, pyridoxine and pantothenic acid are the nutrients most damaged by the sterilization process. Varying results have been described for vitamin losses (5-80% for thiamine; 71-73% for niacin; 49-50% for riboflavin) [28].

Histamine

The histamine compounds are formed from decarboxylation of the amino acid, histidine, through exogenous decarboxylase enzymes that produced by the microorganisms related with fish or environment [29]. Veciana-Nogues, et al. [29] determined that the contents of histamine (HI) (Micrograms per Gram) throughout Canning Process were 0.32, 0.55, 0.40, 0.54 and 0.63 for raw fish, before cooking, after cooking, after packing, and end product respectively. The contents of tyramine (TY) (Micrograms per Gram) throughout Canning Process were 0.32, 0.08, 0.24, 0.17 and 0.15 for raw fish, before cooking, after cooking, after packing, and end product respectively [30]. Histamine poisoning caused by the consuming of fish contains high concentrations of histamine in their flesh [31]. FDA has lowered the histamine defect action level from 100 to 50íg/g and has recommended the use of other biogenic amines related with fish spoilage evaluation [32]. Selmi, et al. [3] Reported that the histamine concentrations increased significantly during storage period but not exceed the acceptable limits. Although tuna and sardine flesh were slightly affected by the canning process, they remained good sources of w3 andw6 fatty acids

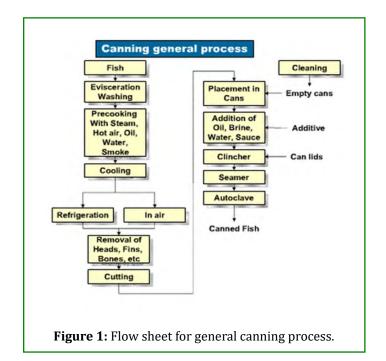
Microbiology

The Egyptian Organization of Standardizations for microbiological aspects of fish products (canned Tuna, canned Sardines, El-Feseekh, salted Sardine, smoked fish and frozen fish), stated that these fish products shouldn't have Clostridium [33-37]. E. coli should be not detecting in Salted, smoked and frozen fish. Coliform bacteria in frozen and smoked fish should be less than 10^3 , 10^1 CFU/g, respectively. TVC should not exceed 10^6 and 10^5 CFU/g in frozen and smoked fish products, respectively. El-Sherif [5] found that there was a highly remarkable decrease in the TPC of the tested canned shrimp samples after canning process, Also, the total bacterial counts of canned shrimp with vegetables were higher than those of canned shrimp with tomato sauce, this can be resulted from the spices and some vegetables contaminated with high. Fatma Arfat [13] reported that canned sardine and mackerel products were found to be free from anaerobic and aerobic thermophilic bacteria during storage at ambient temperature.

Therefore, neither *Clostriduim perfringes* nor *Bacillus cereus* were detected in the canned samples even after 24 months storage. Abd El-Ghafour [15] indicated that the TBC of canned Tilapia was gradual increased as the storage period was prolonged. TBC was increased from 0.113 and 0.107× 10^3 to 0.230 and 0.213 × 10^3 cells/g of steamed canned Tilapia fillets with using either oil or tomato sauce as filling media while was increased from 0.097 and 0.090 × 10^3 to 0.190 and 0.173 × 10^3 cells/g of fried samples with using the same media through storage at ambient temperature.

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A high fungi growth was observed in sun dried and smoked chela (*Laubuka dadiburjori*) after 1 and 2 months of storage compared with canned and frozen chela. This can be from the increasing in water activity and water content [12]. The initial total fungal count of canned chela (*Laubuka dadiburjori*) was 0.95×10^2 (CFU Gg⁻¹), these value increased to 1.13×10^2 and 2.18×10^2 (CFU Gg⁻¹) after 30 and 60 day of storage period respectively. Meanwhile the initial total plate count of canned chela (*Laubuka dadiburjori*) was increased from 1.44×10^4 (CFU Gg⁻¹), to 1.88×10^4 and 2.57×10^4 (CFU Gg⁻¹) after 30 and 60 day of storage period respectively [12,38].



Conclusion

Canned fish products have an economic importance in most countries around the world. During canning process the bacteria and enzymes inactivated by heat treatment, so the canned fish products have a very long shelf life time. Some undesirable effects were occurred during canning process such as loss of essential nutrients, formation of undesirable compounds, browning development and lipid and protein damage which can influence the shelf life of canned products.

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- EOS Egyptian Organization for Standardization (2009) Frozen Fish, Part: 1 Whole Fish, Egyptian Organization for Standardization and Quality, Arab Republic of Egypt, pp: 889-891.
- EOS Egyptian Organization for Standardization (2005e) Smoked Fish, Egyptian Organization for Standardization and Quality, Arab Republic of Egypt, pp: 288.