



Incidence of Some Food Poisoning Microorganisms in Salted fish

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ABSTRACT

A grand total of 90 random samples of salted fish represented by salted sardine, salted *Mugil cephalus* (fesiekh) and salted *Hydrocynus froskahlii* (Mellouha) (30 of each) were collected from different supermarkets at El-Kalyoubia governorate. The collected samples were directly transferred to the laboratory for detection of *Staphylococcus aureus* and *Clostridium perfringens* as food poisoning microorganisms. The obtained results indicated that the mean values of *Staphylococcus aureus* count (cfu/g) in the examined samples of salted sardine, salted *Mugil cephalus* and salted *Hydrocynus froskahlii* were $9.77 \times 10^2 \pm 2.01 \times 10^2$, $4.25 \times 10^2 \pm 0.58 \times 10^2$ and 0/g, with an incidence of 73.33%, 43.33% and 0 % for the same examined samples, respectively. The obtained results revealed also that the mean values of *Clostridium. perfringens* (cfu/g) in the examined samples of salted sardine, salted *Mugil cephalus* and salted *hydrocynus froskahlii* were 2.00×10^2 , $5.53 \times 10^2 \pm 0.7 \times 10^2$ and 0/g, with an incidence of 3.33%, 10% and 0%, for the same examined samples, respectively.

Key words: *Staphylococcus aureus* – *Clostridium perfringens*, Salted fish.

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1. INTRODUCTION

Salted fish products are popular in many countries around the world. Salting is one of the oldest techniques for fish preservation, and essentially intended to increase the shelf-life of the product through depressing water activity by dehydration and salt uptake by the fish muscle. In addition, Sodium chloride is a flavor enhancer as a consequence of its effect on different biochemical mechanisms by reducing or enhancing the enzymatic activity of some enzymes responsible for the development of different organoleptic parameters (Albarracin et al., 2011). However, the current demand for salted fish is driven more by the flavor of the product than for preservation purposes (Ali, Mariyam, 2012). Presence of *Staphylococcus aureus* in a food indicates its contamination from the skin, mouth and/or nose of food handlers. Inadequately cleaned equipment may be a source of

contamination (Thatcher and Clark, 1978). *Staphylococcus aureus* is still a major cause of food poisoning due to ingestion of enterotoxins (Stengel, 1990). The ability to produce such enterotoxin in food is more likely when competing microorganisms were absent (Frazier and Westhoff, 1984). *Clostridium perfringens* spores can reach fish and shellfish in their water habitat (particularly near sewage outfalls), from surfaces of holds in vessels, equipment and utensils used for processing and preparation or from workers. Counts greater than 10^6 cfu/g are necessary to cause illness; such quantities do not reach foods through mere contamination, but accumulate as a result of multiplication of vegetative cells (Bryan, 1980). The present study was carried out to determine the counts of *Staphylococcus aureus* and *Clostridium perfringens* in some marketed salted fish (salted sardine, salted

Mugil cephalus and salted *Hydrocynus Forskalii*).

2. MATERIAL AND METHODS

2.1. Collection of samples:

A total of 90 random samples of salted fish represented by salted sardine, salted *Mugil cephalus* (Fesiekh) and salted *Hydrocynus froskahlii* (mellouha) (30 of each) were collected from different supermarkets at El-Kalyoubia governorate. The samples were directly transferred to the laboratory for count and detection of *Staphylococcus aureus* and *Clostridium perfringens* counts.

2.2. Preparation of samples

It was done according to the technique recommended by ICMSF (1978).

2.3. Determination of *Staphylococcus aureus* count according to ICMSF (1996).

2.4. Determination of *Clostridium perfringens* count was done according to Harmon and Kautter (1978).

3. RESULTS

Table 1 revealed that the mean counts of *Staphylococcus aureus* cfu/g in the examined samples of salted sardine, salted *Mugil cephalus* (fesiekh) and salted *Hydrocynus roskahlii* (Mellouha) were $9.77 \times 10^2 \pm 2.01 \times 10^2$, $4.25 \times 10^2 \pm 0.58 \times 10^2$ and 0/g, respectively. Table 2 showed that there are significant differences between the examined samples of salted fish at ($P < 0.01$) according to their *Staphylococcus aureus* count. Table 3 showed that the mean counts of *Clostridium perfringens* in the examined samples of salted sardine, salted *Mugil Cephalus* (Fesiekh) and salted *Hydrocynus froskahlii* (Mellouha) were 2.00×10^2 , $5.53 \times 10^2 \pm 0.7 \times 10^2$ and 0/g cfu/g respectively. Table 4 revealed that the differences between the examined samples of salted fish were highly significant at ($P < 0.01$) according to the count of *Clostridium perfringens*.

Table (1): Statistical analytical results of *Staphylococcus aureus* count cfu/g in the examined samples of salted fish (n=30).

Salted Fish	+ve samples		Min	Max	Mean±S.E*
	No.	%			
Sardine	22	73.33	1.0×10^2	3.0×10^3	$9.77 \times 10^2 \pm 2.01 \times 10^2$
<i>Mugil. cephalus</i>	13	43.33	1.0×10^2	1.2×10^3	$4.25 \times 10^2 \pm 0.58 \times 10^2$
<i>Hydrocynus froskahlii</i>	0	0	0	0	0

S.E*= Standard error of mean

Table (2): Analysis of variance (ANOVA) of *Staphylococcus aureus* count in the examined samples of salted fish

Source of variance	D.F	S.S	M.S	F. value
Total	89	16183.21		
Between Products (T)	2	2704.95	1352.48	8.73 ⁺⁺
Error	87	13478.26	154.92	

D.F= Degrees of freedom. S.S = Sum squares. M.S = Mean squares. ++ = High significant differences ($P < 0.01$)

Table (3): Statistical analytical results of *Clostridium perfringens* count cfu/g in the examined samples of salted fish (n=30).

Salted Fish	+ve samples		Min	Max	Mean±S.E*
	No.	%			
Sardine	1	3.33	-	-	2.00×10 ²
<i>Mugil cephalus</i>	3	10.00	1.0×10 ²	1.0×10 ³	5.53×10 ² ±0.71×10 ²
<i>Hydrocynus froskahlii</i>	0	0	0	0	0

Table (4): Analysis of variance (ANOVA) of *Clostridium perfringens* count in the examined samples of salted fish

Source of variance	D.F	S.S	M.S	F. value
Total	89	12249.55		
Between Products (T)	2	2995.36	1497.68	14.08 ⁺⁺
Error	87	9254.19	106.37	

D.F= Degrees of freedom. S.S = Sum squares. M.S = Mean squares. ++ = High significant differences ($P<0.01$)

4. DISCUSSION

Results achieved in Table (1) revealed that the mean counts of *Staphylococcus aureus* cfu/g in the examined samples of salted sardine, salted *Mugil cephalus* (fesiekh) and salted *Hydrocynus rosakahlii* (Mellouha) were $9.77 \times 10^2 \pm 2.01 \times 10^2$, $4.25 \times 10^2 \pm 0.58 \times 10^2$ and 0/g, respectively.

There are significant differences between the examined samples of salted fish at ($P<0.01$) according to their *Staphylococcus aureus* count (Table 2). Table (3) showed that the mean counts of *Clostridium perfringens* cfu/g in the examined samples of salted sardine, salted *Mugil Cephalus* (Fesiekh) and salted *Hydrocynus froskahlii* (Mellouha) were 2.00×10^2 , $5.53 \times 10^2 \pm 0.7 \times 10^2$ and 0/g, respectively. The differences between the examined samples of salted fish were highly significant at ($P<0.01$) according to the count of *Clostridium perfringens* (Table 4). The obtained results in the present paper concluded that the examined samples of salted sardine and salted *Mugil cephalus* were more contaminated with *Staphylococcus aureus* and *Clostridium*

perfringens than salted *Hydrocynus froskahlii*. The presence of *Staphylococcus aureus* in The examined salted fish samples suggested the poor personal hygiene of food handlers as well as the organisms may originate from suppurating lesions or from the carrier nostril (Elwi, 1994). The recorded results of *Staphylococcus aureus* count in table (1) were nearly similar to those recorded by Nayel (2007) and Youssef (2011), while Morshdy (1980) and Bashir and Agab (1987) reported higher results. The presence of *Staphylococcus aureus* in salted fish may lead to symptoms of Staphylococcal intoxication which appear within 2-4 hours following consumption of contaminated food (Bergdoll, 1979). Commonly reported symptoms include, nausea, vomiting and less frequently diarrhea, headache, dizziness, with few reported deaths in elderly or very young persons (Varnam, 1991). *Clostridium perfringens* spores can reach fish and shellfish in their water habitat (particularly near sewage outfalls), from surfaces of holds in vessels, equipment and utensils used for processing and preparation or from workers. Greater counts than 10^6 cfu/g are necessary to cause illness, such

quantities do not reach foods by mere contamination, but accumulate as a result of multiplication of vegetative cells (Bryan, 1980). Regarding to the count of *Clostridium perfringens* in, Table (3) indicated that the results of the present paper were lower than those reported by Nayel (2007) who revealed that *Clostridium perfringens* count were detected in 24% of the examined salted sardine samples and 36% of the examined salted *Mugil cephalus* (Fesiekh) samples; Lower results were obtained by Lela-Radwa (2012) who failed to detect *Clostridium perfringens* from all the examined salted samples. In order to control these microorganisms to gain access to salted fish products, and to improve the sanitary status of salted fish processing, periodical medical examination of persons sharing in processing, handling, transporting and salting of fish should be practiced. Licenses should be given to well hygienic fish markets. As well as application and implantation of hazard Analysis and Critical Control Point (HACCP) system in all points of fish manufacturing as a hazardous control system must be applied to ensure maximum safety to consumers.

5. REFERENCES

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