Dept. of Food Hygiene, Faculty of Vet. Med., Assiut University, Head of Dept. Prof. Dr. T.A. El-Bassiony.

MICROBIOLOGICAL QUALITY OF READY TO EAT FRIED FISH

(With Two Tables)

By
Y. HEFNAWY
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الحالة الميكر وبيولوجية للسمك المقلى المعد للإستهلاك الآدمي

يحــــــي حفنـــــاوى

تم جمع خس وعشرون عينة من السمك المقلى المعده للإستهلاك الآدمى من محلات بيسسع الأسماك بأسيوط وحيث تم الفحص الميكروبيولوجى لهذه العينات بالطرق القياسية يهدف تعيين العدد الكلى للميكروبات المعوية وعدد الكور العنقودى اللهبى وعدد الميكروبات السبحية المعوية وبين الفحص الميكروبيولوجى أن العدد الكلى للميكروبات المعوبية وكروبات المعوبية أما العدد الكلى للميكروبات المعوبية وللميكروبات المعوبية فكان يتراوح مابين ١٠ إلى ١٨٥٤ ألى ٥٨٤ الميكروبات المعوبية فكان يتراوح مابين ١٠ إلى ٥٨٤ ١٠ ألى ٥٨٤ ١٠ ألى ٥٨٤ ١٠ ألى ١٤ ألى ١٤ ١٠ ألى ١٤ ١٠ ألى ١٤ ألى ١٤ ١٠ ألى ١٤ ألى ١٤ ألى ١٤ ١٠ ألى ١٤ ١٠ ألى ١٤ ١٠ ألى ١٤ ألى ١٠ ألى ١٤ ألى ١٤ ألى ١٤ ألى ١٤ ألى ١٤ ألى ١٤ ألى ١١ ألى ١٤ أل

SUMMARY

A total of 25 ready to eat fried fish samples were aseptically collected from various food restaurants in Assiut City. These samples were examined bacteriologically by standard procedures for determination of aerobic plate count (APC), and counts of enterobacteriaceae, Staphylococcus aureus and enterococci. The APC ranged from 16x10³ to 188x10¹ colony forming unit (CFU)/gm. The counts of enterobacteriaceae, Staph aureus and enterococci ranged from 10 to 85x10³, 1x10² to 15x10² and 3x10² to 175x10² with a mean values of 4.75x10³, 1.84x10² and 20.1x10² CFU/gm respectively. E.coli, Staph aureus, Clostridium perfringens and Strept faecalis were isolated from the examined samples where 12%, 20%, 8% and 36% of the samples were positive for such organisms respectively. Salmonella and Shigella failed to be recovered from the examined samples. The results of this investigation indicate that foodborne pathogens present in fried fish constitutes a public health hazard.

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INTRODUCTION

Fishes are regarded generally as being more perishable than other high protein foods. Safety in seafood products with reference to bacterial contamination is usually concerned with the possibility of food infection and intoxication. However, food infection and intoxication in seafood products, as other foods, except in instances of scromboid or histamine poisoning, normally is the result of mishandling during and after preparation (NICKELSON and FINNER, 1984).

Literature concerning the microbiology of ready to eat fish is scarce and informative except a report by ELDALY and IBRAHIM (1987) who examined 40 samples of grilled and fried fishes and found that the mean values of aerobic plate count, enterobacteriaceae count, staphylococci count, MPN of coliforms and MPN of E.coli for grilled fish samples were 2x10, 6x10², 6x10³, 2x10 and 2x10² organisms/gm respectively, while the corresponding values for fried fish samples were 9x10 , 2x10², 4x10², 6x10² and 48 organisms/gm respectively. Salmonella and Shigella organisms failed to be recovered from both types of examined fish samples.

For cooked ready-to-eat fishery products, microbiological guidelines for the finished products should include APC, E.coli and Staph aureus. These parameters are useful to evaluate faulty processing and/or handling practices such as inadequate heating, cross contamination with raw products, contamination from workers and inadequate refrigeration which may create hazard (SWARTZENTRUBER, et al. 1980, National Academy of Sciences, 1985).

However, the degree of cooking employed further affects the number and types of organisms. Moreover, organisms normally associated with raw fish are not heat-resistant and are destroyed during heat process. Heat resistant types of organisms may be introduced with spices or other ingredients (NICKELSON and FINNE, 1984).

Furthermore, prepared seafood products can vary greatly in numbers and types of microorganisms because of the addition of nonmarine ingredients as flour, seasonings, nonfat dry milk powder and dried eggs therefore the sourceof contamination may be the ingredients incorporated or the food handlers (KHAN and McCASKEY, 1973; NICKELSON and FINNE, 1984).

No uniform guidelines can be used to interpret the results of bacteriological testing of seafood products where each product must be evaluated on the basis of its own characteristics, and guidelines must be established to practical good manufactuering procedures (NICKELSON and FINNE, 1984).

The purpose of this investigation was to determine the microbiological quality of ready to eat fish and the possible public health hazard associated with its consumption.

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MATERIAL and METHODS

25 samples of ready to eat fried fishes were collected at random from restaurants selling fish in Assiut City. The samples were packaged in sterile plastic bags and brought to the laboratory for microbiological examination.

A 25 gm portions of each sample were blended with 225 ml of 0.1% sterile peptone water in waring blender at high speed for one minute. Serial dilutions from 10 to 10 were made and the bacteriological examinations were done in accordance with the procedures described by BAILEY and SCOTT (1974), CRUICKSHANK, et al. (1975) and SPECK (1984).

All bacteriological media used were of Oxoid unless otherwise specified. Standard plate count agar was used for the aerobic plate count (APC) and plates were incubated at 35°C for 48 hours. Violet red bile glucose agar (VR BGA) was used for total enterobacteriaceeae count (ICMSF, 1978).

For Staphylococcus aureus, appropriate dilutions were streaked on Baird-Parker agar plates (BAIRDPARKER, 1962) and incubated at 35°C for 48 hours. Selected black colonies were tested fro coagulase activity. Enterococcus Selective Differential (ESD) medium (EFTHYMIOU and JOSEPH, 1974) was used for enumeration of enterococci and all magenta colonies were counted as Strept faecalis.

Isolation of Salmonella, Shigella and E.coli was carried out according to the procedures described by ICMSF (1978) and SPECK (1984), whereas the recommended methods outlined by BEERNES, <u>et al.</u> (1980) was followed for isolation of C.perfringens.

RESULTS

Results of enumeration as well as isolation of different organisms from ready to eat fish are shown in Table (1) and (2).

Table (1)
Summerized results of viable counts/gm ready to eat fish

	Minimum	Maximum	Mean
Aerobic plate count	16×10³	118×10 ⁵	22.2x10²
Enterobacteriaceae	10	85×10 ³	4.75×10 ³
Staph aureus	1x10 ²	15×10 ²	1.84×10 ³
Enterococci	3x10 ²	175×10 ²	20.1×10 ²

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Table (2)
Incidence of isolated organisms from examined ready to eat fish samples

Organisms	No. of samples examined	No. positive	Precent positive
E.coli (faecal type)		3-1376	1,377
Staph aureus	25	3	12
C.perfringens	25	5	20
	25	2	8
Strept faecalis	25	9	. 36
Salmonella	25	0	
Shigella	25	0	0

DISCUSSION

The flesh of healthy fish is considered bacteriologically sterile. Fishes may become contaminated during subsequent handling (SHEWAN, 1971). However, coliforms, staphylococci and other mesophilic organisms may be introduced during processing, but there is no evidence that they will multiply unless there is inadequate temperature control (ICMSF, 1980).

Table (1) revealed that the aerobic plate count of the examined fried fish samples ranged from 16x10³ to 118x10 with a mean value of 22.2x10 CFU/gm, while the mean enterobacteriaceae count was 4.75x10³ with a minimum of 10 and a maximum of 85x10³ CFU/gm of examined samples. Our results seemed to be some what higher than that obtained by ELDALY and IBRAHIM (1987) who recorded an average of 9x10 and 2x10² CFU/gm for APC and enterobacteriaceae count respectively.

Regarding staphylococci and enterococci counts they varied from $1x10^2$ to $15x10^2$ and $3x10^2$ to $17x10^2$ with a mean values of $1.84x10^2$ and $20.1x10^2$ CFU/gm respectively.

However, ADESIYUN (1983) reported that the mean staphylococcal count of fried fish in Nigeria was 2.5×10^6 /gm which is higher than the obtained results, but ELDALY and IBRAHIM (1987) recored an average of 4×10^2 /gm fried fish samples.

SURKIEWICZ et al. (1968) found that fried breaded fish had APC values of less than 2.5×10^{-4} /gm, MPN of colfiorm had less than 10/gm, and no more than 10% of the units were positive for E. coli or staphylococci.

Seafoods are susceptible to all of the common food poisoning organisms. Staph aureus, Salmonella, Shigella and C. perfringens have occasionally been responsible for foodborne disease outbreaks from consumption of fishery products (NICKELSON and FINNE, 1984).

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The summarized results presented in Table (2) pointed out that E. coli, C. perfrings, Staph aureus and Strept faeclis were recovered from 12%, 8%, 20% and 36% of the examined samples respectively, while Salmonella and Shigella failed to be detected in this investigation.

Safety in sea food products with reference to bacterial contamination is usually concerned with the possibility of foodborne infection and intoxication. The present study proved that ready to the fried fish is considered as a public health hazard due to post cooking contamination and the results agree with the finding of NICKELSON and FINNE (1984) that food poisoning in sea food products is the result of mishandling during and after preparation.

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