INCIDENCE OF SOME ENTEROTOXIGENIC FOOD POISONING MICROORGANISMS IN CHICKEN MEAT PRODUCTS

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SUMMARY

Forty - five packaged random samples of cooked and half cooked chicken meat products represented by chicken shawarma, fillet and wings (15 for each) were collected from different retail markets in Cairo and Giza. The incidence of S. aureus, E. coli and C. perfringens in fillet was 26.66%, 20%, 26.66%, respectively, while equals 0.0%, 13.33%, 20%, in shawarma, respectively. The incidence in wings reaches to 13.33%, 20%, 6.66%, respectively. Six number of S.aureus strains were isolated from fillet and wings examined for enterotoxin production and revealed that only one strain isolated from wings had the ability to produce enterotoxin types A, B, C and D (multiproducer). While S.aureus failed to be isolated from shawarma. Concerning C. perfringens 5 toxigenic strains classified as 3 C.perfringens type A and 2 C. perfringens type D, while 3 strains were non- toxigenic. For E. coli the serological typing revealed 8 untypable strains which were not enterotoxigenic.

INTRODUCTION

Chickens occupy the major role in production and consumption among poultry in Egypt. Chicken meat becomes the second most popular type of meat eaten after red meat. Chicken meat is characterized by ease during preparation consistent quality and the availability of the wide range of pre-packed, branded, raw, ready to eat and serve products (Shedeed, 1999).

Bacterial agents are incriminated in foodborne infection and intoxication outbreaks in industrial and developing countries, which increase day by day (Stevenson and Bernard, 1995).

Study of the different causative bacterial agents of food poisoning is one of the pioneering efforts to improve poultry industry with the resultant protein gap covering. Determination of types of some microorganisms in most carcasses is important from the stand point of public health for judging effectiveness of sanitary handling during processing (Butler el al., 1979).

The food poisoning organisms of most concern with meat such as enteropathogenic *E. coli* and *C. perfringens* are associated with enteric contamination (ICMSF, 1980), moreover, *C. perfringens* is a very common cause of human foodborne diseases and in most outbreak cases the food involved are cooked meat or poultry products containing high levels of viable cells (Labbe, 1989; Labbe and Harman, 1992).

Beside E. coli and C. perfringens, S. aureus continues to be a major cause of foodborne intoxication and its presence in food constitutes an important hygienic problem for food processors, food service, workers and consumers (Bergdoll, 1989).

The possible sources of *E. coli* and other pathogens contaminated the ready to eat meat products included inadequate sanitary practices, inadequate heat treatment and the presence of pathogens on different surfaces continuously or occasionally contaminating the finished products (Gibbons et al., 2006).

Thus, this study was conducted to insure in what range we need to apply and control the hygienic measures on chicken meat products along its processing and retailing.

MATERIAL AND METHODS

Samples: A total of 45 random samples of chicken meat products of cooked shawarma and half-cooked fillet and wings packaged units (15 for each) purchased from different retail markets in Giza and Cairo. The packaged of samples transfered in an ice box without delay and under hygienic conditions to the laboratory for the bacteriological examination.

* Preparation of the samples (APHA, 1992):

Twenty five grams of food sample were homogenized with 225 ml of sterile buffered peptone water (0.1%) to give a dilution of 1/10. One ml of the clear homogenate was mixed with 9 ml of buffered peptone water (0.1%), then decimal dilutions were prepared.

* Estimation of Escherichia coli count (MPN):

The most probable number was conducted as recommended by (FAO, 1992), isolation and biochemical identification according to Feng et al. (1998) and serologically according to Cruickshank et al. (1975).

* Staphylococcus aureus count:

The procedure of FAO (1992) was followed.

* Enumeration of Clostridia organisms:

Enumeration of Clostridia organisms was con-

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fucted according to ICMSF (1978) and Bergeyís Manual (1986).

* Isolation of C. perfringens:

The method was applied according to ICMSF (1978) and identification according to Buchanan and Gibbons (1975).

- * Demonstration of *C. perfringens* toxin by dermonecrotic test (Sterne and Batty, 1975).
- * Detection of S. aureus enterotoxins:

It was done according to Donelly et al. (1967); Oda et al. (1979) and Shingaki et al. (1981) using the SET-RPLA kit for the detection of staphylococcal enterotoxins A, B, C and D.

- * Detection of Enterotoxins of E. coli strains: Preparation of E. coli strains for recovery of enterotoxin according to Evans et al. (1973) and Alderate and Robertson (1977).
- * Detection of heat labile enterotoxin (LT.) among E. coli isolates:

 Serologically using Oxoid VET-RPLA kit for detection of E. coli heat labile enterotoxin.
- * Detection of heat stable enterotoxin (ST) among E. coli isolates according method by Giannella (1976).

RESULTS

Table (1): Statistical analytical results of isolated microorganisms.

samples		Fi	llet*		74.	Shaw	arma* *		II.	Wir	igs*	
Microorganisms	Min	Max	Mean	SD	Min	Max	Mean	SD	Min	Max	Mean	SD
S cureus	1x10 ²	1.1x10 ³	6x10 ²	2.l x10		•	•		2x10 ²	5x10 ²		
E coli	2.8x10	1.5x10 ²	7.4x10	1.3x10	7	2.3x10	• 1	197	4	75	28.5	4.6
Clostridia	8x10 ²	7.6x10 ⁴	17.5x10 ²	1.6x10 ²	4x10 ²	3x10 ³	8.66x10 ²	8.lx10	1.5x10 ²	3.9x10 ³	17.3x10 ²	3.9x10

* Half-cooked

** Cooked

Data represented as mean ± SD

Table (2): Incidence of isolated microorganisms from chicken meat products (n = 15).

samples	E	Fillet	Shaw	Shawarma	Wings	ıgs
/						•
Microorganisms	No.	%	No.	%	No.	%
S. aureus	4	26.66	0	0	2	13.33
E. coli	·	20.0	,2	13.33	သ	20.0
C. perfringens	4	26.66	သ	20.0	1	6.66

Table (3): Enterotoxin types of isolated S. aureus.

Chicken	No. of isolated	Enterotoxigenic	Types of
samples	strains	strains	enterotoxins
Fillet*	4	1	
Shawarma* *		-	•
Wings*	2	1	A,B,C,D

Table (4): Typing of C. perfringens strains (n = 15).

Chicken	No. of	T	Toxigenic strains	strain	S	Non-toxigenic	xigenic
samples	isolated	1		н	J	str	strains
	strains	No.	%	No. %	%	No.	%
Fillet*	4	-	25	_	25	2	50
Shawarma**	3	2	66.6	•	ı	1	33.3
Wings*	_	1	•	Н	100	•	·

* Half-cooked

** Cooked

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piscussion

The revealed results gave a profile about the hygienic and microbiological status of some cooked (shawarma) and half-cooked (fillet and wings) chicken products and showed that these products could harbour the food poisoning microorganisms easily and the achieved results must give more attention to follow up the hygienic rules in the processing, handling and storage of such products.

Table (1) showed that the mean count levels of S. aureus was $6 \times 10^2 \pm 2.1 \times 10$ in fillet, while not detected in shawarma and wings. *E. coli* recorded mean count value/g using (MPN) equals 7.4 x 10 \pm 1.3 x 10 and 28.5 \pm 4.6 in each of fillet and wings, respectively. While, the mean count level of clostridia organisms reached to 17.5 x $10^2 \pm 1.6 \times 10^2$, 8.66 x $10^2 \pm 8.1 \times 10$ and 17.3 x $10^2 \pm 3.9 \times 10$ in fillet, shawarma and wings, respectively.

In this regard, Hafez et al. (1987) detected *S. aureus* in chicken meat product (burger) in mean level count of 4.6 x 10³ CFU/g, Hefnawy and Moustafa (1990) detected *S. aureus* 1 x 10² in ready to eat chicken products, while Hashim (2003) detected 5 x 10² CFU/g. Shabana and Ouf (2003) found *S. aureus* in chicken shawarma in range mean count of 2.72 - 3.36 log count. Ahmed (2004), Gad (2004), Essa et al. (2004) detected *S. aureus* mean count reached 6.77 x 10³,

 3.8×10^3 , 3.8×10^3 and 1.37×10^4 CFU/g from chicken nuggets, wings, fillet and luncheon, respectively.

For *E.coli* mean count level, Soriano et al. (2000) found *E. coli* from 3 - 2.400 MPN/g, Shabana and Ouf (2003) detected *E. coli* from chicken shawarma in range of 1.94-2.11 log count and Hashim (2003) found coliform 2.9 x 10² in chicken meat product. On the other hand, Warburton et al. (1988) failed to detected *E.coli* from processed poultry products.

Concerning clostridia organisms Osman (2005) detected clostridia in a mean count ranged from 2.3 x 10² - 3.3 x 10³ CFU/g. while, *C. perfringens* in count mean range 3 x 10² - 9 x 10³ CFU. On the other hand, Wen and McClane (2004) failed to detect *C. perfringens* in chicken product samples.

As shown in Table (2) the incidence of S. aureus, E. coli and C. perfringens were 26.66%, 20.0% and 26.66% in fillet samples, while in shawarma was 0.0%, 13.33% and 20.0%, respectively. In wings samples the incidence represented in 13.33%, 20% and 6.66% for S. aureus, E. coli and C.perfringens. In this respect, many researchers studied and isolated food poisoning microorganisms in different chicken meat products as; Bryan (1988) found S. aureus (8%), and C. perfringens (5.2%); Hefnawy and Mosutafa (1990) isolated S. aureus and E. coli at rate of

and E. coli (8.8%). High incidence (73.6%) repractices during processing, storage and retailing such pathogens indicated substandard hygiene all products with conclusion that the presence of coliforms and S. aureus were particularly high in in all meat products and added that the levels of ed faecal coliforms, S. aureus and C. perfringens poultry and meat product samples and he detectthe other hand, Murugkar et al. (1993) studied taminate the flesh (Avens and Miller, 1970). On flesh of healthy live poultry is sterile but bacteria while, Nasr et al. (2007) isolated C. perfringens Osman (2005) isolated C. perfringens in range average of 37% from meat and poultry samples, Miwa et al. (1998) found C. perfringens in an products. Regarding, C. perfringens incidence found E. wings, respectively. While, Sharma et al. (2005) isolated E. coli (20%) and (12%) from fillet and 21.4% and E. coli (7.1-35.7%). Ahmed (2004) Ouf (2003) isolated S. aureus from shawarma in 10%, respectively from hot wings. Shabana and lated S. aureus and E. coli in rate of 18% and (7.5%) and E. coli (6.3%). Shaltout (2002) isothe other hand, Girgis (2002) recorded S. aureus corded by Chang et al. (2001) for S.aureus. On (2000) found the incidence of S.aureus (7.6%) 20% and 10%, respectively. While, Soriano et al on the skin surface such as S. aureus may conranging between 8.3 to 14%. In this respect, the 10-50% from different chicken meat products, coli (14%) from meat and chicken

The results in table (3) illustrated that 2 strains of S. aureus isolated from wings samples were examined against enterotoxins production and typing, as a result one strain only (50%) considered enterotoxigenic and could produce enterotoxin A, B, C and D types (multiproducer), while we can't detect any toxigenic S. aureus strains from fillet. As previously recorded in the results that shawarma is negative for presence of S. aureus and that may be due to exposure of these samples to efficient heat treatment (cooked) and avoid of recontamination after processing and packaging.

chicken meat and the same author found that the enterotoxin type C was predominated. Staphyloenterotoxigenic S. aureus at rate of 29.64% from ture than by bacterial competition (Herten et al., S. aureus grow better but toxin production ap-(Anon, 1986). In the absence of competitors gestion of enterotoxin produced during growth of types of foodborne illness and results from the incoccal food poisoning is one of the most common from chicken meat. Chang et al. (2001) isolated of S. aureus strains were enterotoxigenic isolated terotoxigenic, while, Zaki (1998) found 54.16% isolated S. aureus strains from chicken were en-In this respect Adesiyun (1984) found 36% of the peared to be influenced more by growth temperaenterotoxigenic strains of S. aureus

The direct extraction of examined chicken meat samples (fillet, wings and shawarma) and typing

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for presence of S. aureus enterotoxins showed negative results (not detected). Sally Rose et al. (1989) reported that the detection of preformed enterotoxins (SETs) is therefore important in epidemiological studies outbreaks food poisoning and in routine quality appraisal monitoring of foods during manufacture. On the other hand, Wieneke et al. (1993) provided that enterotoxins were detected in foods in the absence of viable S. aureus and most contamination took place at home followed by restaurant and shops.

S. aureus on chicken cuts insured that there is potential incidence reached to 43.3% (Sliva et al., 2002). Colombari et al. (2007) stated that strains of S. aureus isolated from food and food handlers implicated as the etiologic agent of an outbreak of staphylococcal food poisoning involving 180 people occurred in Brazil, in April, 1998.

Regarding table (4) the results showed the toxins typing of *C. perfringens* strains by *I/D* inoculation in Guinea pigs as follows in fillet 4 strains isolated were classified as 2 strains toxigenic, one type A (25%) and the second (25%) type D and the other two strains (50%) were non toxogenic. Three strains isolated from shawarma and classified as 2 (66.6%) type A and one (33.3%) was non-toxigenic. Only one strain of *C. perfringens* type D was isolated from wings.

In this respect, Osman (1999) reported that out of 22 strains of *C. perfringens* 17 of them provided

enterotoxigenic type A, which is usually involved in food poisoning outbreaks and noticed that all C. perfringens recovered from cooked products proved toxigenic and the non-toxigenic were only allocated to the frozen raw products.

ted through poultry (Mulder, 1997) and (Dalton of outbreaks of human food poisoning transmitcal reports incriminated C.perfringens as a source (Abigail and Dixie, 1994), and the epidemiologistrains of C. perfringens causes pronounced diarpossess the enterotoxigenicity. The toxigenic high, and not all strains of these microorganisms tries where consumption of meat and poultry is ing cause of bacterial foodborne illness in coun-Labbe (2003) stated that C. perfringens is a lead-A and 12.9% were type D. In this regard, Lin and ic C. perfringens strains and typing as 70.8% type while, Nasr et al. (2007) detected 83.9% toxigenfound 23% of the strains provided to be toxigenic terotoxigenicity of C. perfringens. Osman (2005) By examination of chicken meat products for enet al., 2004). 12 hrs after consumption of contaminated food rhea and abdominal cramps generally appeared 8-

To prevent *C. perfringens* infections cooked beef and poultry meat should be refrigerated promptly and reheated thoroughly (internal temperature 75°C) before serving (Enan, 2006).

Examination of *E.coli* isolated strains (8 strains) for serological identification, the results revealed

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(Foster, 1987). heat stable toxin (ST). From the public health thermolabile toxin (LT), while 4 only produced enterotoxigenicity, and found none produced ic diarrhea in children, food poisoning outbreaks of gastroenteritis, acute infantile diarrhea, sporadpoint the serotyped E. coli causes human cases ed 45 strains of E. coli must be free from E.coli according to E.S. types (Klipstein et al., 1978), and cooked sample that the properity of enterotoxigenicity of E. coli labile (LT) and heat stable (ST) yielding negative (2000). On the other hand, Sosa et al. (1988) testis clearly not restricted to enterotoxigenic seroresults (non-toxigenic). It is benefit to mention ity of isolated E. coli strains for detection of heat untypable strains, in addition the enterotoxigenicisolated from food for

It could be concluded that the potentially of some enterotoxigenic microorganisms (S. aureus, E. coli and C. perfringens) that may be present or harbour the chicken meat products, so that barriers could be put at the processing lines to minimize the likelihood of such organisms surviving or multiplying to levels which may limit these products acceptability.

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