

Bacteriological and chemical evaluation of some heat treated chicken products.

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ABSTRACT

A total of sixty commercially produced coated poultry products samples (heat treated) representing, 20 each of (chicken breast, nuggets and wings) were examined to evaluate its quality in terms of bacterial and chemical attributes. Bacteriological analysis showed that the mean values of *aerobic mesophilic*, *Psychrotrophic, staphylococcal* and *coliform* counts were $21.6 \times 10^3 \pm 3 \times 10^3$, $11.5 \times 10^3 \pm 2.2 \times 10^3$, $20 \times 10^2 \pm 11 \times 10^1$ and 3.33 ± 0.6 in chicken breast, $42.2 \times 10^3 \pm 3.3 \times 10^3$, $20.7 \times 10^3 \pm 4.5 \times 10^3$, $14 \times 10^2 \pm 4 \times 10^2$ and 3.33 ± 0.8 in nuggets and $60.4 \times 10^3 \pm 10.4 \times 10^3$, $38.5 \times 10^3 \pm 5.4 \times 10^3$, $10 \times 10^1 \pm 7$ and 79.6 ± 6.2 in wings, respectively. Meanwhile chemical analysis showed that, the mean values of TVN and TBA were 27.4 ± 2.6 and 0.59 ± 0.04 for breasts samples, 28.4 ± 1.13 and 0.48 ± 0.12 for nuggets and 27.0 ± 2.1 and 0.65 ± 0.14 for wings. When the obtained results compared with the Egyptian Standards, the examined breast samples were found to be of better quality than the other types of chicken product samples, Meanwhile the incidence of rejected samples as it contain coagulase positive *Staph. aureus* were 20, 30 and 20 % respectively.

Keywords: bacteriology- chemical - (heat treated) chicken meat products.

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1.INTRODUCTION

Commercialization of heat treated chicken products has increased in the recent decades because of its practicality and convenience to prepare, and increase of consumer power (Suderman, 1983; Smith et al., 1985 and Sunderland, 1992). A part from the nutritional aspect, the growing tendency to spend less time on food preparation has lead to a great demand for timesaving heat treated frozen food products also they are very attractive food items due to its low production cost (Antonova et al., 2003 and Patton, 2005) moreover, they are favored by consumers because of the increased palatability provided by a soft and moist interior juiciness and tenderness. Production of heat treated chicken meat products is a complex procedure involving particle size reduction, blending forming, coating and cooking. The possibilities existing in each of these steps adds to the potential variety of these products but also increases the potential for problems if done incorrectly (Sams, 2001). Poultry meat and their products often get contamination from different sources starting from defeathering, evisceration and subsequent handling during processing in plant (Levin, et al., 2001 and Houf, et al 2002). Many efforts were done to produce a product free from pathogens of public health hazard and with low microbial count improving its keeping quality and keeps its nutritive value to be safe and of high quality. However, many other problems exist like contamination during cutting or maceration of tissues and lose of nutritive values. During freezing of chicken meat products, the growth of many types of microorganism will cease while others especially psychrotrophic bacteria

can grow until the medium freezes (Davis and Board, 1998). Psychrotrophic bacteria are responsible for many undesirable changes in flavor, odor, texture and color of the food products. The presence of Staphylococci and, in particular Staph.aureus in the retail breaded chicken products is a potential health risk for consumers since the pH and aw values of these kinds of products are favorable for Staph.aureus growth. The thermal process used during their manufacture can limit staphylococcal contamination but cannot eliminate preformed toxins (Pepe et al., 2006). So the present study was planned to examine some heat treated products for its quality and safety for human consumption through assessment for bacteriological quality in addition, chemical analysis will be performed to assure quality in the aspects of consumer acceptability, degree of freshness

2. MATERIALS AND METHODS

2.1. Samples

A total of 60 samples of coated heat treated chicken products (20 samples of each) breasts, nuggets and wings were collected from different super markets. All samples were subjected to bacteriological and chemical examination.

2.2. Bacteriological examination

Preparation of samples Homogenate: Ten grams from each sample were homogenized with 90 ml sterile 0.1 % peptone solution in a sterile polyethylene bag for 1.5 minutes using stomacher (Lab-blender 400). One ml from the sample original homogenate was added to a test tube containing 9 ml 0.1% sterile peptone water to provide a dilution of 10^2 . Similarly a tenfold serial dilution was prepared (APHA, 2001). And the following bacteriological investigations were performed.

2.3. Enumeration of Total aerobic bacterial count (APHA, 2001)

One ml from each dilution transferred with sterile pipette to each of two separate sterile

Petri-dishes, and then about 10 ml of the sterile standard plate count agar melted at 45°C were poured to each Petri-dish. Inoculated plates after being mixed and solidified were incubated at 37°C for 24-48 hours. Colonies were recorded and counted as total bacterial count cfu/g.

2.4. Enumeration of total Psychrotrophic count

Method applied according to (APHA, 2001), the inoculated plates are incubated in inverted position at 7°C for 10 days. Accordingly, the total *psychrotrophic* bacterial count per gram was calculated on plates containing from 30 to 300 colonies.

2.5. Enumeration of Total Staphylococcal Count

From each dilution 0.1 ml was spread onto a dry surface of double sets of Baird parker agar plate (Oxoid CM 275, SR 54). Inoculated plates were incubated at 37°C for 48hours. Typical colonies of *Staph. aureus* (black shining convex colonies, 1-1.5 mm in diameter with narrow white margin and surrounded by a clear zone extending into opaque medium) were enumerated and the average number per gram was calculated (APHA, 2001) The purified *Staph. aureus* isolates were identified through different biochemical tests, catalase test, coagulase test (tube test) (Quinn, et al., 2002)

2.6. Enumeration of Total Coliform (MPN/g) (APHA, 2001)

Estimation of *coliforms* was done by using most probable number technique with MacConkey's broth tubes. A series of 3 fermentation tubes containing MacConkey;s broth and inverted Durham's tubes were inoculated with 1 ml from the previously prepared 10th fold serial dilutions. After thorough mixing, inoculated and control tubes were incubated at 37 °C 24-48 hours. Tubes showing acid and gas were considered as positive for the test. From the laboratory records, the most probable number (MPN) of *coliforms /g*. was calculated by matching with (MPN) table.

2.7. Chemical analysis

All samples were examined for the following chemical analysis to estimate their compatibility with the Egyptian

3. RESULTS

standards (EOSQC, 2005). Determination of basic nitrogen (Total Volatile Nitrogen) (TVN) and thiobarbituric acid (TBA) values were estimated as an indicator of the degree of products freshness. Samples will be analyzed in accordance to the methods of FAO, (1980).

Table (1): Mean values of bacteriological examination in examined chicken meat samples (Breasts, Nuggets and wings). (N: 20)*

Samples	T. aerobic mesophilic count	T. Psychrotrophic count	T. Staph. Count	T. Coliform count (MPN/g)	
Breast	$21.6 \times 10^3 \pm 3 \times 10^3$	11.5×10 ³ ±2.2×10 ³	$20 \times 10^{2} \pm 11 \times 10^{1}$	3.33±0.6	
Nuggets	$42.2 \times 10^3 \pm 3.3 \times 10^3$	$20.7 \times 10^{3} \pm 4.5 \times 10^{3}$	$14 \times 10^{2} \pm 4 \times 10^{2}$	3.33±0.8	
Wings	$60.4 \times 10^3 \pm 10.4 \times 10^3$	$38.5 \times 10^3 \pm 5.4 \times 10^3$	$10 \times 10^{1} \pm 7$	79.6±6.2	
* N [.] Number of examined samples					

* N: Number of examined samples

Table (2): Acceptability of chick	ten meat products samples	according to EOSQC (2005)
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Samples							
Brea	Breasts		ets	Wings		Stan danda	Microorga
	Unaccepted samples				Standards	nisms	
N/20	%	N/20	%	N/20	%		
0	0	3	15	0	0	< 100 MPN/g	Coliform
4	20	6	30	4	20	Free	Staph. aureus

Table (3): Mean values of Chemical analysis of (heat treated) chicken meat products and their acceptability according to *EOSQC*, (2005).

Samples	TVN		TVN mg/100 g	TBA	TBA < 0.9 mg/kg	
		N/20	%		N/20	%
Breasts	27.4±2.6	4	20	0.59±0.04	0	0
Nuggets	28.4±1.1	6	30	0.48±0.12	4	20
Wings	27.0±2.1	4	20	0.65±0.14	4	20

4. DISCUSSION

Bacteriological examination of chicken meat products clearly indicated that the chicken wings samples had significantly higher bacterial load than either breast or nuggets, the results in table (1) showed that the mean aerobic mesophilic plate count (cfu/g) of examined breast, nuggets and wings was $21.6 \times 10^3 \pm 3 \times 10^3$, 42.2×10^3 $\pm 3.3 \times 10^3$ and $60.4 \times 10^3 \pm 10.4 \times 10^3$ nearly similar results was observed in ELHoti. (2011) while Osman, (1997; 2001) and Sofroni et al., (2008) recorded higher mesophilic counts for frozen chicken products. The aerobic plate count gives an idea about the hygienic measures applied during processing and also helps in the determination of the keeping quality of the product. The highly aerobic count indicates contamination of raw material or unsatisfactory processing as well as it may unsuitable environmental be due to condition during storage. Results recorded in table (1) for *psychrotrophic* count were $11.5 \times 10^3 \pm 2.2 \times 10^3$, $20.7 \times 10^3 \pm 4.5 \times 10^3$ and $38.5 \times 10^3 \pm 5.4 \times 10^3$ this increase of the mean values in wings samples may be due to the prolonged storage period rather than the breast and nuggets samples or due to the fluctuations in storage temperatures. Higher values were observed in EL-Shora, (1990) who found that the log mean values of APC and total psychrotrophic counts are 7.43 and 6.78 for frozen chicken products and Abd EL-Magied, Walaa, et al., (2009) who found the *psychrotrophic* count was $1.43 \times 10^5 \pm 0.37 \times 10^5$ /g in breast samples and $4.28 \times 10^6 \pm 0.38 \times 10^6$ /g in wings. In contrast lower values were reported by Zaki-Nadia (1994). In general. the contamination of chicken meat products with great number of psychrotrophs could be attributed to the neglected sanitary measures adapted during intensive preparation, processing, handling and packaging as well as cold storage (Cenci et al., 1990). Furthermore, the contaminated equipments and knives are probably the principle contributing factors to high

psychrotrophic counts of such chicken meat products (Davies and Board, 1998). Data shown in table (1) revealed that the mean values of staph. aureus count in examined samples was $20 \times 10^2 \pm 11 \times 10^1$, 14×10^2 $\pm 4 \times 10^2$ and $10 \times 10^1 \pm 7$ respectively, lower findings were observed in AL-Dughavm and Altabari (2010); ELHoti, (2011) and Wang et al., (1976) this high count of staphylococcal sp. Indicate bacterial contamination during packing and handling by the workers. The mean *coliform* count in chicken breast, nuggets and wings was 3.33 ± 0.08 $3.33 \pm .06$ and 79.6±6.2 (MPN/g), as illustrated in table (1) it seems to be low due to the effect of freezing which minimize the count of *coliform*. This is in accordance to Hamada, (2012) and James, et al., (1992). Generally, the presence of coliform in chicken meat products is considered as an indicator for improper handling and unhygienic conditions after slaughtering, de-feathering, and washing fresh chicken carcasses. Our results were agreed with Frazier and Westhoff (1983) and Hashim (2003) while Abdel-Haffeiz, (1999) could not detect it from nuggets. According to the legal requirements of Egyptian Organization for Standardization and Quality Control (EOSQC, 2005), as shown in table (2), it is evident that 20, 30 and 20 % of breast, nuggets and wings samples respectively had Staph. aureus above the permissible limit. Presence of Staph. aureus may be attributed to inadequate heat treatment, unhygienic handling practices, use of dirty containers. faulty storage and transportation, so the hands and clothes of employees in the production of chicken meat should be over looked (Duffrenne et al., 2001) nearly similar results recorded by Pepe el al., (2006). It was clearly evident from the obtained results that only nuggets samples (15%) had *coliform* count above the EOSOC, (2005), while all samples of both breast and wings were accepted. So Staph. aureus count considered as an index of the sanitary quality of examined samples. From

a food safety perspective, it is recognized

that *Staph. aureus* is an enterotoxinproducing pathogen but that the concentration needs to exceed 10^5 cfu/ml for sufficient toxin to be produced to cause human illness (Hill,1983).

Results achieved in table (3) indicated that the mean values of TVN mg\100g were 27.4±2.6, 28.4±1.13 and 27.0±2.1 and the mean values of TBA were 0.59 ± 0.04 , 0.48 ± 0.12 and 0.65 ± 0.14 for breast, nuggets and wings respectively. Also, according to Egyptian Organization for Standardization (EOSQC, 2005) for heat treated chicken meat, the percentage of unaccepted samples of our products under investigations was 20, 30 and 20 for TVN analysis and 0, 20 and 20 % for TBA analysis in breast, nuggets and wings sample respectively. Regarding the examined samples, nearly similar results were recorded by Shams El-Din and Ibrahim (1990) and Shedeed (1999).

TVN can be considered as a reliable indicative measure for the quality of various food articles specially chicken and chicken cuts-up. In general, TVN in chicken cuts-up may be increased as the days of storage increased (Reddy et al., 1970). Our results were agreed with those obtained by Hassanin-Fatin and Hassan (2003). Regarding the examined samples, nearly similar results were recorded by Shams El-Din and Ibrahim (1990) and Shedeed (1999).

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التقييم البكتيري والكيميائي لبعض منتجات الدواجن المعاملة حراريا

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الملخص العربى

تواجه صناعه منتجات الدواجن المعامله حراريا كثير من المشاكل الصحية اثناء التصنيع مما يؤثر بصوره كبيره على صحة المستهلك ولتحديد جوده هذه المنتجات تم جمع عدد عشرين عينه من كل من صدور ، قطع النجتس واجنحه الفراخ الجاهزه المطهي لتقييم جودتها بكتريولوجيا وكيميائيا وقد اظهرت النتائج ان متوسط العدد الكلي البكتيري والبكتريا المحبه للبروده ومكروب العنقودي الذهبي والميكروبات القولونيه هو 103×3 ±01×10, 210×10 ×103 ×2.2 ±103×10 ×101×11 ×101×10 ×102 ±201×20 ±201×10 ×101×10 ×102 ±201×20, 103×10 ±201×20, 103×10 ±201×20, 103×10 ±201×20, 103×10 ±201×20, 103×10 ±201×20, 103×10 ±201×20, 103×10 ±201×10 ±200 ±30.00 ±30.00 ±30.00 ±30.00 ±30.00 ±30.00 ±30.00 ±30.00 ±20

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