

Animal Health Research Institute
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INCIDENCE OF *LISTERIA MONOCYTOGENES* IN SOME MEAT PRODUCTS AND POULTRY

(With 2 Tables)

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(Received at 12/12/1998)

مدى تواجد الليستيريا مونوسيتوجين فى بعض منتجات اللحوم والدواجن

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يعتبر مرض الليستيريويزيس من أهم الأمراض التى تصيب الإنسان والذي يسببه ميكروب الليستيريا مونوسيتوجين وخاصة عند تناول الأطعمة التى لاتتعرض لدرجة حرارة كافية وبسبب خطورة هذا الميكروب على صحة المستهلك أجريت هذه الدراسة لمعرفة مدى تلوث بعض منتجات اللحوم والدواجن بهذا الميكروب ، لذا تم جمع عدد ١٥٠ عينة (٣٠ عينة لكل من اللحم المفروم والسجق وقطع الدجاج و ٤٠ عينة من اللنشون و ٢٠ عينة من العينات الجاهزة للأكل بواقع ١٠ عينات من كل من (الدجاج المشوى على الفحم ، الكفتة المشوية على الفحم) أمكن عزل ميكروب الليستيريا مونوسيتوجين من ٩ (٣٠%) من اللحم المفروم و ٤ (١٣,٣%) من السجق و ١٤ (٤٦,٦%) من الدواجن بينما لم يتم عزل الميكروب من ٤٠ عينة من اللنشون, كما أمكن عزل عترة واحدة (١٠%) من الدجاج المشوى على الفحم بينما لم يتم عزل الميكروب من ١٠ عينات من الكفتة بعد الشوى .

SUMMARY

Listeria monocytogenes has been recognized as major foodborn pathogen especially foods which are not exposed to heat treatment. So a total of 150 samples of raw and cooked ready-to eat meat meals collected from different supermarkets and restaurants were examined for the presence of *L. monocytogenes*. Out of 30 samples of each raw ground beef, sausage and raw poultry, the positive samples were 9(30%), 4(13.3%) and 14(46.6), respectively. While the organism could not be detected in 40 samples of luncheon. In cooked ready to eat poultry meat the organism

was detected in one sample only (10%) and failed to be detected in 10 samples of grilled kofta. The source of product contamination, thermal and preservative used and public health significance were discussed.

Keywords: *Listeria Monocytogenes*, Meat Poultry

INTRODUCTION

Listeria monocytogenes has become of significant concern in recent years as a food borne pathogen (El-Kest and Marth, 1988 and Gellin *et al.*, 1991).

Meat and meat products have frequently been found to be contaminated with *L. monocytogenes* and may serve as vehicles of this pathogenic bacterium (Elis Cherova *et al.*, 1979; Nicholas, 1985; Breuer and Prandl, 1988; Johnson *et al.*, 1988; Karches and Teufel, 1988 and Schmidt *et al.*, 1988). The frequent occurrence of *L. monocytogenes* and its ability to survive and proliferate at refrigeration temperatures provokes accumulation of this pathogen in contaminated meat products during cold storage and hence presents of potential risk for consumers (Doyle, 1988).

In humans the primary manifestations of listeriosis included septicemia, endocarditis, pneumonia, conjunctivitis, pharyngitis, cutaneous papules and pustules, urethritis and meningitis (Banwart, 1989).

As *L. monocytogenes* is an ubiquitous organism, it can be shed from the intestine of both man and animals without any apparent clinical symptoms (Gracey and Collins, 1992). Therefore the presence of the organism during the processing of meat and meat products becomes unavoidable (American Meat Institute, 1988 and Lowry and Tiong, 1988). Also the post processing contamination of ready to eat meat products with *Listeria* spp. could be of potential hazard (Johnson *et al.*, 1988).

Although *L. monocytogenes* has been primarily associated with dairy products it is associated also with other food of animal origin used for human consumption and has been isolated from ground beef (Nicholas, 1985 and Brakett, 1988). Nicholas and Vidaud (1987) found *L. monocytogenes* in 26% of samples of frozen minced meat samples whereas the incidence in ground beef varied from 28 % as recovered by Skovgaard and Morgen (1988) and to 58% as found by Truscott and

McNad (1988). Also, Farber *et al.* (1989 a) found that 38 of 44 (86.4 %) ground meat and 6 of 30 (20%) sausages samples contained the organism. Furthermore Fathi and Saad (1994) found that 1 of 7 (14.20 %) of sausage and zero in luncheon were contaminated by *L. monocytogenes*. Breer and Schopfer (1988) isolated *L. monocytogenes* from 18% of poultry while Jennifer *et al.* (1990) detected the organism in 25.6 % to 60 % of meat and meat products according to places of samples.

Therefore the present study was designed to survey the most popular meat products and poultry for the presence of *L. monocytogenes*.

MATERIAL and METHODS

(1) Collection of samples:

A total of 150 random samples of meat products including frozen ground beef, sausage, and raw poultry, (30 samples each) and 40 samples of luncheon taken from different supermarkets, as well as 20 samples of ready to eat poultry and Kofta after grilling on coal fire (10 samples each) were taken from various restaurants in Assiut City. Each sample was wrapped separately and aseptically in sterile polyethylene bag, then labelled and transferred as quickly as possible to be tested bacteriologically for the presence of *L. monocytogenes*.

(2) bacteriological analysis:

25 grams of each sample were homogenized with Moulinex type blender equipped with metallic flask in 225 ml of Listeria enrichment broth (Oxoid CM 862 and Oxoid SR 141) for 1 min and incubated at 30°C for 48 h. After incubation one loopful was subcultured on Listeria selective agar (Oxoid CM 856 and Oxoid SR 140). Colonies of Listeria were visible after 24 h. at 37°C and the incubation was continued for a further 24 h. to detect slow growing strains according to Oxoid Manual (1990).

Identification of the isolated strains was tested according to Seeliger and Jones (1986) and Loyett (1988). Colonies were picked on tryptone soya agar incubated at 30°C for 24 h. and confirmed through Gram stain reaction, catalase, motility at 21°C in motility agar medium and Kilgar iron agar for glucose and lactose fermentation, hydrogen sulphide production, Beta haemolysis on defibrinated sheep blood agar at 37°C for 48 h. and carbohydrate fermentation (Mannitol) and nitrate reduction.

Mouse pathogenecity was performed to confirm *L. monocytogenes* according to Seeliger and Jones (1986).

RESULTS

The obtained results were tabulated in Table 1 and 2.

Table 1: Incidence of *L. monocytogenes* in some products meat and Poultry.

Meat product	No. of examined Samples	Positive samples	
		No.	%
Ground beef	30	9	30
Sausage	30	4	13.3
Luncheon	40	-	-
Raw poultry	30	14	46.6

Table 2: Incidence of *L. monocytogenes* in ready to eat Poultry and Kofta

Cooked ready to eat meats	No. of examined samples	Positive samples	
		No.	%
Grilled Poultry	10	1	10
Grilled Kofta	10	0	0

DISCUSSION

The results in Table (1) revealed that *L.monocytogenes* could be detected in 9(30%) of ground beef, 4(13.3%) sausage and 14(46.6%) of raw poultry, while none of the analyzed luncheon samples was found to contain *L. monocytogenes*.

Ground beef:

These results are nearly similar to those of (McClain and Lee, 1988; Schwartz *et al.*, 1988; Skovgaard and Morgen 1988; WHO, 1988; Grau and Vanderlinde, 1992 and Elgazzar and Sallam (1997) where they could detect *L. monocytogenes* in arround (30-50%) of raw ground meat. On the other hand, high percentage of ground meat positive samples (86%) was detected by Farber *et al.* (1989 b) and Lee and McClain (1989) (77.3%).

The reason of such variation may be due to hygienic measures and differences in methodology.

Chicken:

The obtained high results (46.6%) lie within the range recorded by Gitter (1976), McClain and Lee (1988), Pini and Gilbert (1988) and Skovgaard and Morgen (1988) who reported the presence of the organism from 14.7 % to 60% of raw poultry. On the other hand Farber *et al.* (1989) which could isolate the organism from 56 % of chicken legs. The higher incidence of *L. monocytogenes* in poultry meat may be due to poultry slaughter practices which are responsible for some of this contamination and the frequent occurrence of *L. monocytogenes* in bird feces also, it has been speculated that deep litter of poultry houses may be the source (Skovgaard and Morgen, 1988).

Sausages:

The results in Table (1) show that (13.3%) of samples were positive. These results were nearly similar to those of Johnston *et al.* (1988) who found that 5 of 42 samples (contained *L. monocytogenes* 11.9 %). Fathi and Saad (1992) who found that 1(14.28 %) of fermented sausage were positive for *L. monocytogenes* after drying. Sausage fermentation and drying process can be regarded as means of reducing but not eliminating listeria from Sausage (Glass and Doyle, 1989 a). Higher results were recorded by Farebe *et al.* (1989a) who found that 6 of 30 (20%) fermented sausage contained the organism. The lowest percentage of positive samples may be due to the high susceptibility of *L. monocytogenes* to combined chemical preservatives added to sausage (Shahamat *et al.* 1980, El-Shenawy and Marth 1988, and Wederquist *et al.*, 1994).

Luncheon:

L. monocytogenes could not be detected in luncheon samples under investigation. Such result was similar to that obtained by (Fathi and Saad (1992). On the contrary, Wilson (1989) and Furrer *et al.* (1991) revealed the presence of such organism in 4 and 6% of the examined Luncheon samples respectively. However, the lowest percentage may be due to spices and temperature during manufacture and good hygiene.

Cooked ready to eat samples:

In ready to eat poultry, *L. monocytogenes* could be detected in 1(10%) and could not be detected in grilled Kofta (Table 2). Anon (1989) and Coote *et al.* (1991) found that 70°C for 2 min. cause reduction in numbers of *L. monocytogenes*. While Karaio annoglou and

Xenos (1981) could isolate *L. monocytogenes* from meat balls grilled for 15 min. at 78 - 85°C internal temperature. Kilduff (1987 a,b) and Wilson (1989) could isolate *L. monocytogenes* from about 2% (2/100) of corned beef samples. These variations may be due to that the organism could survive cooking or the poultry were contaminated after cooking (Boyle et al. 1990).

Results of monitoring program of ready to eat products in U.S.A. revealed that 5 - 12% of products sampled in (1987) and 10-13 % in 1988 were contaminated with *L. monocytogenes* (1989). Several studies occurred by Glass and Doyle (1989 b) indicated that post processing contamination by *L. monocytogenes* may be a hazard.

From the above show it is important to give sufficient time during cooking, and good hand washing for food workers, good cleaning and sanitizing of the plant and elimination of product residue, using of unconventional meat product additives such as sodium lactate, nisin which act against the growth of *L. monocytogenes*. Good manufacturing practices should decrease the incidence of Listeria.

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