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A SURVEY OF SOME SELECTED FOOD ITEMS FOR THE PRESENCE OF LISTERIA MONOCYTOGENES AND OTHER LISTERIA SPECIES

(With 2 Tables)

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بعض الأغنية التي لا تتعرض لحرارة كافية في إعدادها لها دور خطير في نقل كثير من الأمراض ، من هذه الأمراض مرض Listeriosis والذي يببه ميكروب له من هذه الأمراض مرض Listeriosis والذي يببه ميكروب المدروب Lamonocytogenes من المدروب المدر

SUMMARY

Various retail food products were analyzed for the presence of listeria species, including 10 luncheon, 7 sausage, 9 burger, 100 kareish and 50 damietta cheese samples. Out of 7 sausage samples, one (14.20%) was positive for Listeria monocytogenes, while the organism failed detection in the examined lucheon and burger samples, and it was found in one sample of kareish (1%) and one of damietta cheese (2%), respectively. However, Listeria innocua could be isolated from one (10%), 2(28.57%), 3(3%) and one (2%) of the examined luncheon, sausage, kareish and damietta cheese samples, respectively. Listeria welshimeri could be detected in one sample of burger (11.1%) and one sample of kareish cheese (1%). The public health importance of Listeria species and the suggestive measures were recommended.

FATHI & NAGAH

INTRODUCTION

Listeria monocytogenes is a foodborne pathogen which has lately been a cause of great concern not only for the food industry but also for regulatory agencies as well FARBER and LOSOS (1988) and WHO (1988). Listeria monocytogenes is widely distributed in the environment, it has been isolated from soil, water and animals and occurs widely in various fresh foods such as raw milk and meats (BRACKETT, 1988).

Listeria monocytogenes is associated with meningoencephalitis, septicemia. pneu-monia, endocarditis, urethritis and abortion (GRAY and KILLINGER, 1966).

The pathogen has been isolated from food processing environment (COX et al., 1989) and finished food products (ANONYMOUS, 1987).

The World Health Organization, in its recommendations to the food industry regarding Listeria, stressed the importance of gathering information about the ability of various food to support growth of the pathogen (WHO, 1988).

Meat and dairy products as well as varieties of other foods, the processing of which does not include a final listeriocidal heat treatment are now widely recognized as possible sources of foodborne listeriosis (JAMES et al., 1983 and SCHLECH et al., 1983). Although L. monocytogenes has been primarily associated with dairy products, it is associated also with food animals used for human consumption (BRACKETT, 1988) and has been isolated from ground beef (NICHOLAS, 1985). In 1987, the Food Safety Inspection Service initiated a survey of meat products to determine the extent of contamination at the retail level. NICHOLAS and VIDAUD (1978) found L. monocytogenes in 26% of samples of frozen minced meat, whereas the incidence in ground beef varied from 28% (SKOVGAARD and MORGEN, 1988) to 58% (TRUSCOTT and McNAB, 1988). FARBER et al. (1989) found that 38 of 44(86.4%) ground meats and 6 of 30(20%) of sausages contained the organism.

The presence of L. monocytogenes in milk products has been documented. It has been isolated from various milk products in the market place. The pathogen behaves differently in different kinds of cheese, it survived for more than one year in CHEDDAR, 140 days in Colby, 90 days in Feta, and grow in Camembert chesse (EL-GAZZAR and MARTH, 1991).

RODLER and KORBLER (1989) isolated L. monocytogenes from 2% of examined cheese samples. HIRD (1987) stated that L. monocytogenes was recovered from a sour cream like product and cottage cheese by-products. Listeria ivanovii is another pathogenic species usually associated with sheep. In addition, a variety of Listeria species has been found in meat and meat products. Isolation of L. innocua from mea is common and the incidence of this organism is higher than the incidence of L. monocytogenes (BREER & SCHOPFER, 1980 and BREUER & PRANDL, 1988). Listeria

FOOD LISTERIA MONOCYTOGENES, OTHER LISTERIA

seeligeri and i. welshimeri have been documented recently to cause infections in human (ROCURT et al., 1986 and ANDRE & GENICOT, 1987).

The use of Listeria species other than L. monocytogenes as indicators of the presence of that organism has been proposed WHO (1988).

Therefore, this investigation is to survey some food products for the presence of L. monocytogenes and other Listeria species.

MATERIAL and METHODS

Food samples which included 10 luncheon, 9 burger, 7 sausage, 50 damietta and 100 kareish cheese were collected from retail local markets and supermarkets and examined for the presence of Listeria monocytogenes and other Listeria species.

Enrichment procedure:

10 g of each food product were placed aseptically in an enrichment broth consisting of Tryptose broth containing 40 Ug/ml Nalidixic acid and 30 u/ml Polymyxin B sulphate as described by HOFER (1983). The broth was incubated at 37°C for 24 h.

Isolation and identification of Listeria species:

After incubation a loopful of enrichment broth was streaked on the surface of Tryptose agar supplemented by 40 Ug/ml Nalidixic acid and 30 u/ml Polymyxin B sulphate and incubated at 37°C for 24 h. Colonies resemling those of Listeria were Gram stained. Confirmatory tests were done on each isoalte suspected to be L. monocytogenes as described by SEELIGER (1961) and LOVETT (1988). Mouse pathogenicity testing was performed to confirm L. monocytogenes according to SEELIGER and JONES

RESULTS

The obtained results were recorded in Tables (1 & 2).

DISCUSSION

The results in Table (1) revealed that L. monocytogenes could be detected in 1(14.28%) of sausage, 1(1%) kareish and 1(2%) damietta cheese samples, while non of the analyzed luncheon and burger samples were found to contain Listeria monocytogenes. These results are nearly similar to those of JOHNSTON et al. (1988) who

Assiut Vet.Med.J., Vol. 27, No. 54, July 1992.

FATHI & NAGAH

found that 5 of 42 samples 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 Listeriae from sausage (GLASS and DOYLE, 1989).

Higher results were recorded by FARBER et al. (1989) who showed that 6 of 30(20%) fermented sausage contained the organism.

Concerning cheese samples, similar results were obtained by RODLER and KORBLER (1989) and GARAYZABAL and GENIGEORGIS (1990). L. monocytogenes has been reported to survive during manufacture and store of some cheese varieties (STAJNER et al., 1979).

As shown in Table (2), all Listeria isolates from food samples were identified as L. monocytogenes, L. innocua, and L. welshimeri. It is clear that 1(10%) of luncheon, 2(28.57%) sausage, 3(3%) kareish cheese and 1(2%) damietta cheese samples were positive for L. innocua. Literia innocua was previously isolated from meat, and its incidence is higher than the incidence of L. monocytogenes (BREER AND SCHOPFER, 1988).

However L. welshimeri could be isoalted from 1(11.1%) burger and 1(1%) kareish cheese samples. VANDERLINE and GRAU (1991) recorded higher results. L. innocua and L. monocytogenes were previously isolated from beef roast by SKOVGAARD and MORGEN (1988).

The use of any Listeria species as an indicator of the presence of L. monocytogenes has been proposed (JOHNSTON et al., 1990).

In the meantime, control of listeriosis could be achieved by n awareness of the ubiquity of the organism and especially of those environments that favour multiplication. Thus, the colaboration of medical, veterinary and food bacteriologists; farmers, food manufacturers, and technologists is essential.

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FOOD LISTERIA MONOCYTOGENES, OTHER LISTERIA

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FATHI & NAGAH

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FOOD LISTERIA MONOCYTOGENES, OTHER LISTERIA

Table (1): Isolation of Listeria monocytogenes from some food produts

Type of samples	No. of examined samples		Positive samples		
			No.	%	
Luncheon	10		0	0	
Sausage	7		1	14.28	
Burger	9		0	0	
Kareish cheese	100		1	1 .	
Damietta cheese	50		1	2	

Table (2): Isolation of Listeria species from some food produts

	No. of examine	L. monoc	No. of position L. monocytogenes		samples innocua	L.welshimeri	
		NO.	%	No.		No.	%
Luncheon	10	0	0	. 1	10	0	0
Sausage	7	1	14.28	2	28.5	7 0	0
Burger	9	0	0	0	0	1	11.1
Kareish cheese	100	. 1	. 1	- 3	3	1	1
Damietta chees	e 50	1	2	- 1	2	0	0