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Prevalence of Listeria monocytogenes in Human in Dhamar Governorate/Yemen

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ABSTRACT: This research was conducted to investigate the existence and spread of Listeria monocrtogenes among different ages of human. Three hundred and ten samples (310) were collected (100 male blood, 110 female blood, and 100 Placenta). The results revealed that the overall prevalence of L. monocytogenes in total samples was N 61 (19.7%), the isolation percentage from the Female blood N 22 (20.0%) was higher than from the Male blood N 18 (18.0%), while the isolation percentage from the placenta reached to N 21 (21.0%).

When we study the isolation rate of L. monocytogenes from Male blood according to Directorates , the results showed that the high rate of isolation were N 11 (31.4%) in Thamar Directorate, following Gahran N 2 (15.4%), next Ans N3 (10.7%), then Alhada N2 (9.1%), but there is no sample was isolate in Anes Directorate. Whereas the rate of isolation in Female blood according to Directorate were N 10 (29.4%) in Thamar , following N 2 (20.0%) in Anes , N 5 (17.9%) in Alhada, N 2 (14.3%) in Gahran , and N 3 (12.5%) in Ans Directorate . The highest isolation rate of the bacteria agents from placenta were in Gahran N4 (36.4%), then N9 (33.3%) in Alhada, N5 (26.3%) in Ans, N1 (16.7%) in Anes, and finally Thamar Directorate N 2 (5.4%). The results indicated that there are significant differences at (p<0.05) in Prevalence of L.monocytogenes in Placenta according to Directorates .

This study showed that the high prevalence rate of L. monocytogenes from Male group aged more than (60) years, had higher opportunities to be infected with the microbe than other age groups, in this age group the isolation percentage was (30.0%), followed by the Male between forty one to fifty years was 3 (27.3%), then from thirty one to forty years where 3 (16.7%), next that from Male between the ages of twenty one to thirty years were 3 (12.5%), finally less than twenty years where 3 (11.1%).

Whereas the higher percentage of isolated L. monocytogenes from Female according to ages were (27.3%) between (21-30) years, followed by female with ages between forty one to fifty years, more than fifty one years, thirty one to forty years, less than twenty years, their infection percent were 3 (25.0%), 2 (22.2%), 3 (15.8%) and 2 (7.7%) respectively, and results illustrated that the highest rate of infection among pregnant women were in age groups less than 20 years with the percentage of (29.4%), followed the ages between 31-40 years were 9 (27.3%), then between 21-30 years where 7 (14.6%), while not isolated from the age between (41- 50) years. The relationship between months and prevalence of L. monocytogenes during period of study indicated that the highest isolation rate occurred in August (44.4%), and September (34.3%).

From this study we concluded that the prevalence of L. monocytogenes in human in Dhamar Governorate was high, and that human infection with these bacteria can occur at any stage of life, especially the period of age after 60 years in males and the stages in which the rate of pregnancy increases in females.

Due to the importance of this study we recommend further researches and studies on L. monocytogenes in different governorates of Yemen . Also working on creating specialized a national center deal with further studies and diagnosis of L. monocytogenes , and emphasis for inter the diagnosis of this bacteria within the routine work of the laboratories in hospitals. This study also recommends the dissemination of health awareness through the media, audio and visual media and all categories of the community male and female as well. Highlighting the health risks resulting from infection with these bacteria, and follow the scientific standards should be considered and adopted in the field of public health to prevent their transmission to humans.

Key words: Listeria monocytogenes, Human, Dhamar Governorate, Yemen.

INTRODUCTION

The genus Listeria is a member of the family Listeriaceae , and currently includes 17 species: L. aquatica, L. booriae, L. cornellensis, L. fleischmannii, L. floridensis, L. grandensis, L. grayi, L. innocua, L. ivanovii, L. marthii, L. monocytogenes, L. newyorkensis, L. riparia, L. rocourtiae, L. seeligeri, L. weihenstephanensis, and L. welshimeri¹, of which L. monocytogenes poses the greatest threat to human health²⁻³.

Listeria monocytogenesis a virulent strain causing febrile gastroenteritis in healthy people , and invasive diseases in vulnerable populations such as pregnant women , newborns , the very young , the elderly and people who are immunocompromised , the incubation period range between 1 to 90 days , with a medium of 3 weeks , and the median incubation period in noninvasive listeriosis is one day with a range of 6 hours to 10 days $^{4-6}$.

Most cases of listeriosis are reported in the third trimester, but the women may become infected with L. monocytogenes (L. monocytogenes) at any time during pregnancy. Usually three to seven days after the onset of symptoms, a woman may abort the fetus or have premature delivery. listeriosis may result in spontaneous abortion in the first trimester. While in case of later stages of pregnancy, the result may be stillbirth or birth of a critically sick newborn ⁷. The critical target sub population for L. monocytogenes infection are the Pregnant women, and the pregnancy has been associated with about 18 - fold increased risk of developing disease than the non pregnant female population ⁸.

L. monocytogenes is a Gram - positive, rod-shaped bacterium that form single short chains, nonspore forming bacterium which measures 0.4 μ m in width and 1 to 1.5 μ m in length, and it is a facultative intracellularanaerobe. The bacterium is motile being flagellated, especially below 33°C via 1-5 peritrichous flagella; it can be resistant to the effects of freezing, drying, and heat. L. monocytogenes has the ability to grow at low temperatures; thus, allowing it to grow in refrigerated foods with optimum temperature between 30-37°C⁹⁻¹⁰.

Nilsson¹¹ stated that the L. monocytogenes was a ubiquitous saprophytic bacterium and capable of causing a severe infection in humans. The organism has been recognized for 84 years, known as a human pathogen for approximately 80 years, and a food borne etiology was confirmed 27 years ago.

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L. monocytogenes is a zoonotic facultative intracellular pathogen that can invade into and multiply inside of host cells. Infection with L. monocytogenes in humans can cause two forms of listeriosis, the non – invasive gastrointestinal form which develops as a typical febrile gastroenteritis and invasive form which can also be acquired by the fetus from its infected mother via the placenta. After L. monocytogenes enter into the host across the gastrointestinal epithelium, it disseminates via the blood and the lymph stream, then the bacteria establish in the spleen and the liver. From here the L. monocytogenes can be re-released into the blood stream and subsequently cross the fetoplacental barrier in pregnant women, besides the blood-brainbarriers for entry into the CNS, where they can cause meningitis and meningoencephalitis¹²⁻¹³.

Listeriosis is a Listeria-related illness characterized by flu- like symptoms; fever is generally present in patients with bacteremia, other nonspecific symptoms such as malaise, fatigue, muscle aches, gastrointestinal symptoms may also occur. When the infection with L. monocytogenes spreads to the nervous system, symptoms in this case may progress to include severe headache, confusion, loss of balance or convulsions and stiff neck ¹⁴⁻¹⁵.

In Yemen the prevalence and the pathogenic role of L. monocytogenesin Humans have been neglected, therefore the objectives of this research were to study the prevalence of L. monocytogenes among different ages of human in Dhamar Governorate and to determine the relationship between months and prevalence of L. monocytogenes in human during the period of study. Also the study of the epidemiological data on L. monocytogenes helps in the establishing public health action that could halt transmission and therefore acquisition of the infection and aid the therapeutic program to eradicate the bacterium.

MATERIALS AND METHODS

1- Study Design and Sampling:

Three hundred and ten (310) human samples (100 Male blood , 110 Female blood, and 100 Placenta) were collected from General Thamar Hospital and some private medical laboratories. The samples were collected in dry sterile container according to (20), Personal information about patients with samples recorded; include Age, Sex, Place of residence and Directorates.

2- Isolation and Identification of L. monocytogenes:

In laboratory, the isolation of L. monocytogenes from blood was done under sterile conditions by placed 10 ml fresh blood into blood culture bottles containing 20 ml of Listeria Enrichment Broth (LEB) and incubated at 37°C for 48 hours, subcultures were then made on Listeria Oxford Medium Base (OXA) and on Brain Heart Infusion Agar (BHIA) plates by streaking, inoculated plates were incubated at 37°C for 48 hours in microaerophilic atmosphere. The isolation of L. monocytogenes from placenta was done by taken 5x5 cm from placenta and put in sterile Morter, added peptone water and grinded after cut into

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small pieces by sterile scissors , after that transfer samples into special container containing 20 ml of Listeria Enrichment Broth (LEB) and incubated at 37°C for 48 hours , lop full from (LEB) streaking on Listeria Oxford Medium Base (OXA) and Trypticase Soy Agar (TSA) , then incubated the plates at 37°C for 48 hours in microaerophilic atmosphere ¹⁶⁻¹⁷.

After incubation period, all plates from blood and from placenta were examined to determine the properties of typical colonies of L. monocytogenes. These colonies are stained with Gram stain. Isolates are small, smooth and appear pale blue-green when viewed from the side (45 angle) with a beam of white light, gram positive with exposure the smear to Crystal violate for 1 min, slightly curved, tiny rods with rounded ends, often occurring in pairs at an acute angle ¹⁸, but old cultural may appear gram negative¹⁹. The identification of L.monocytogenes was confirmed by Biochemical tests as listed in (Table 1)²⁰.

3- Statistical Analysis

Statistical Analysis was done by using the Statistical Package for Social Sciences (SPSS, version 20). Used for descriptive statistics the crosstabs and used for possibility taste the Chi-Square.

RESULTS

From 310 samples of different types of human blood and placenta, 61 (19.7%)) were gave a positive result for isolation of L. monocytogenes from total human samples. These results include 18 (18.0%) positive samples from Male blood, 22 (20.0%) positive samples from Female blood, and 21 (21.0%) positive samples from placenta (Table 1 and Figure 1). There were no significant differences between prevalence of L. monocytogenesin Human (Male and Female) in (p > 0.05).

When we study the prevalence of L. monocytogenesin Human blood according to Sex, we noticed that Female is more exposed (20.0%) to infection withL. monocytogenes, compared with Male infection rate (18.0%) (Table 3, Figure 2).

The prevalence of L. monocytogenes in male according to Directorate was high in Thamar Directorate(31.4%), following Gahran (15.4%), Ans (10.7%), then Alhada (9.1%), but L. monocytogenes did not isolate from male blood in Anes Directorate (Table 4 and Figure 3). Also the rate of isolation in Female blood according to Directorate were (29.4%) in Thamar Directorate, following (20.0%) in Anes, (17.9%) in Alhada, (14.3%) in Gahran, and (12.5%) in Anes Directorate(Table 5 and Figure 4). From 100 samples of placenta, the high rate of isolation were in Gahran (36.4%), then (33.3%) in Alhada, (26.3%) in Ans, (16.7%) in Anes and finally Thamar Directorate (5.4%) (Table 6, Figure 5). The results indicated that there are significant differences at (p<0.05) in Prevalence of L. monocytogenesin Placenta according to Directorates

This study showed that the isolation rate of L. monocytogenes from Male group aged \geq 60 years were (30.0%) , followed by the group with age between 41- 50 years (27.3%), from 31– 40 years (16.7%), then from 21 – 30 years (12.5%), finally in the group < 20 years showed (11.1%) (Table 7 and Figure

6).When we study the prevalence of L. monocytogenes in female according to age , we found that the age of Female between 21– 30 years of 12 participants (27.3%) have a chance to be infected more than the other age groups, followed by female with ages between 41– 50 years, \geq 51years, 31 – 40 years ,less than 20 years , their infection percent were; 3 (25.0%), 2 (22.2%), 3 (15.8%) and 2 (7.7%) respectively (Table 8 and Figure 7).

Other consideration which influenced the infection with L. monocytogenesin Pregnancy woman may be the ages factor, isolation of L. monocytogenes from woman less than 20 years were 5 (29.4%), followed the ages between 31-40 years were 9 (27.3%), then between 21- 30 years were 7 (14.6%), while we did not found L. monocytogenes in the bigger ages 41–50 years, as shown in (Table 9, Figure 8).

When we study the relationship between months and prevalence of L. monocytogenes in total human samples during the study period, the highest isolation rate in August 16 (44.4%), September 12 (34.3%), and April 12 (33.3%) (Table 10, Figure 9).

The relationship between directorates and abortion cases in General Thamar Hospital from 2009 to 2012 according to statistics recorded in Thamar Health Bauru were illustrated in (Table 11, Figure 10).

DISCUSSION

Listeriosis is a serious invasive illness caused by the bacterium L. monocytogenes, which primarily afflicts pregnant women, neonates, very young, older adults, and people with weakened immune systems.

Listeria monocytogenes is a zoonotic facultative intracellular, Gram-positive flagellated food borne pathogen causing serious disease worldwide. L. monocytogenes infection in humans can result in septicemia, encephalitis, meningitis, abortion, premature birth, and stillbirth. Mortality due to listeriosis is very serious leading to death in about 30% of the human cases , while in vulnerable populations such as fetuses, infants, elderly, and immunocompromised individuals , the mortality rate may be reach to 75% .

This bacterium are widely distributed in the environment , with ability to survive and multiply under low temperatures and have high tolerance ranges to both salt concentrations (10%) as well as a broad range of pH . Domestic and wild mammals, birds, and man may be asymptomatic carriers of Listeria in their intestinal flora. About 40 mammalian species have been shown to harbor Listeria, which can be isolated from intestinal samples of up to 10% of the human population.

In the study at hand, Three hundred and ten samples(310) were collected included 100 male blood, 110 female blood, and 100 Placenta samples. The overall prevalence of L. monocytogenes in total human samples was N 61 (19.7%), as shown in (Table 2, Figure 1). There were no significant differences between prevalence of L. monocytogenes in Human (Male and Female) in (p > 0.05). Our result was approach with percentage found by ²¹, whom found the prevalence of L. monocytogenes infection in human in Finland, at rate (23.0%). AlsoFugett²² in New York reported that the prevalence of L. monocytogenes in known in the prevalence of L. monocytogenes in L. monocytogenes in Known in the prevalence of L. monocytogenes in further the prevalence of L. monocytogenes in further the prevalence of L. monocytogenes in Known in the prevalence of L. monocytogenes in further the prevalence of L. monocytogenes in Known in the prevalence of L. monocytogenes in Known in the prevalence of L. monocytogenes in further the prevalence of L. monocytogenes in Known in the prevalence of L. monocytogenes in the prevalence of L.

monocytogenes was 21 of 24 (87.5%) among people consuming the food contaminated with L. monocytogenes.

Our result incompatible with the study conducted by ²⁴ in Argentine, who found that the patients with Listeriosis represented (46.72%) of 122 cases. Also our result are disagreement with ²⁵ who found that the prevalence of L. monocytogenes in human were (72.2%) in Italy. In another hand, ²⁶ mentioned that the prevalence of L. monocytogenes among human samples were (7.3%), and ⁶ in Iran, observed that (8.8%) out of 125 human samples were positive for L. monocytogenes.

Schuppler and loessner²⁷ mentioned that the organisms are well adapted to the conditions in the gastrointestinal tract and pursue different strategies to counteract changes in acidity, osmolality, oxygen tension, or the challenging effects of antimicrobial peptides and bile. The finding that the bacteria are able to colonize and persist in the gall bladder and suggest the occurrence of long-term and chronic infections and demonstrates the ability of pathogenic Listeria to survive within the various microenvironments of the gastrointestinal tract.

The total prevalence of L. monocytogenes in human samples according to Sex were 40 (19.0%), and the isolation rate in female blood 22 (20.0%) were higher than in Male blood 18 (18.0%), as show in (Table 3 and Figure 2).

Our result were compatible with²⁸ in Switzerland, whom isolated L. monocytogenes from blood in rate (21%), also they obtained that (21%) of the cases were of bacteremia, (40%) of meningitis, and (39%) of meningoencephalitis. Many studies were inconsistent with our finding , Lukinmaaet al.,²¹ in Finland , found that L. monocytogenes were detected in (57%) of male samples and (42%) of female samples. Lake et al.,²³ in New Zealand, mentioned that the isolation rate of L. monocytogenes from male and female were (51.3%) and (48.7%) respectively. Whereas in Tehran²⁹ did not found L. monocytogenes in a total of (398) blood samples. However, Leong et al.,³⁰ declare that L. mOnocytogenes has the ability to cross the epithelial barrier of the intestinal tract to cause more serious infection throughout the body including bacteremia, it can also cross the blood-tissue barrier which allows the bacteria to infect organs such as the brain or uterus , where it can cause severe life-threatening infections such as meningitis , encephalitis , spontaneous abortion , or miscarriage.

Table 4 and Figure 3; shown that the prevalence of L. monocytogenes in male blood according to Directorates. The prevalence of L. monocytogenes in Thamar Directorate were high (31.4%), following Gahran Directorate (15.4%), then Ans (10.7%), and (9.1%) in Alhada Directorate, while we did not isolate L. monocytogenes from Anes Directorate. The results indicated that there were no significant differences at (p > 0.05).

Also Table 5 and Figure 4, show that the prevalence of L. monocytogenes in female blood according to Directorates. The prevalence of L. monocytogenes were (29.4%), (20.0%), (17.9%), (14.3%)

and (12.5%), in Thamar, Anes, Alhada, Gahran and Ans Directorate respectively. The results indicated that there were no significant differences at (p>0.05).

The reason of high numbers of infection cases in Thamar Directorate maybe due to the presence of hospitals and health center in this urban directorate, and the population in this directorates were more awareness about the importance of diagnosis of the disease, as well as the high level of education in this directorates. In contrast, poor hygiene and sanitation and the close proximity to animals in rural directorates all contribute to easy and frequent acquisition of any enteric pathogen, including listeria.

As shown in Table 6 and Figure 5, the prevalence rate of L. monocytogenes in placenta according to directorates, were (36.4%) in Gahran , followed by Alhada (33.3%), Ans (26.3%), Anes (16.7%), and finally Thamar Directorate (5.4%).

The reason of high percentages of prevalence of L. monocytogenes in Gahran Directorate may be due to the low level of education in this rural Directorate, and lack of health awareness, as well as the absence of health program for the pregnancy women. The results indicated that there were significant differences at (p<0.05).

From Table 7 and Figure 6, the prevalence rate of L. monocytogenes in Male blood of different age's groups. The prevalence of L. monocytogenes in the age more than 60 years were 6 (30.0%), followed by the Male with age between 41 to 50 years were 3 (27.3%), then from 31 to 40 years were 3 (16.7%), after that from 21 to 30 years were 3 (12.5%), finally the age less than 20 years were 3 (11.1%). There were no significant differences at (p>0.05) in prevalence of L. monocytogenes in male blood according to Age.

Our results were in agreement with²⁸, who found that L. monocytogenes was isolates from (42%) of the patients had an underlying disease and (54%) were > 65 years of age. Patients with bacteremia were significantly older than those with meningitis or meningoencephalitis (median ages,75,69, and 55 years, respectively).

In Denmark Larsen et al.,³¹ determined L. monocytogenes in Male between 52 to 82 years old with septicaemia rate (3.70%), and 64 years old with meningitis were (7.14%), finally the bigger age were 86 years old with septicaemia (16.67%).Our result non-agreement with study conducted in New Zealand by²³ where the prevalence rates of L. monocytogenes were (45.2%) among the peoples aged over 60 years, while the prevalence rate (34.3%) among the peoples over 80 years.

When we compared between prevalence of L. monocytogenes in female blood according to age Table 8 and Figure 7, the result showed that the age group between 21 to 30 years was the highest age of infection 12 (27.3%), followed by age group between 41 to 50 years 3(25.0%), age group more than 51 years 2 (22.2%), then age group between 31 to 40 years 3 (15.8%), and finally the age group less than 20years 2 (7.7%). The results indicated that there were no significant differences at(p>0.05) in Prevalence

of L. monocytogenes in female blood according to Age. In Atlanta, ³² prevalence of L. monocytogenes infection was (31%) in age group < 65 years, and (53%) in patients aged \geq 65 years.

In Denmark ³³ studied the prevalence of L. monocytogenes infection between different age groups. The prevalence of L. monocytogenes infection were (0.4%), (3.7%), (7.3%), (12.1%) and (22.0%) for the age groups (0-59), (60-69), (70-79), (80-89) and (90+) years respectively. Other study in Denmark³¹, found different rates between ages 58, 63, 76 and 80 years, the rates were (50.0%), (80.0%),(26.7%) and (20.67%) respectively.

In London Gillespie et al.,³⁴ reported that the prevalence rate of L. monocytogenes in patient < 60 years were (33%) out of 385 cases, while the prevalence rate in patient \geq 60 years were (66%) out of 783 cases. Current results were non-agreement with result found in New Zealand²³, where they found (54.8%) of the L. monocytogenes infection with age 60 years, and (65.7%) with age 80 years.

According to age groups in pregnancy woman the highest isolation rate of L. monocytogenes were in age less than 20 years (29.4%), followed by the woman with age between 31 to 40 years (27.3%), then from 21 to 30 years (14.6%), while we did not isolate L. monocytogenes from women in age group between 41 to 50 years, this finding can be explained in (Table 9, Figure 8). The results indicated that there were no significant differences at (p > 0.05) in Prevalence of L. monocytogenes in Pregnancy woman according to age in Thamar province.

Women infected during pregnancy may pass L. monocytogenes to the fetus, either transplacentally or at birth. Infection in a fetus may result in stillbirth or preterm delivery while infection in a neonate may present as meningitis or septicemia. Rare outbreaks in neonatal nurseries have been attributed to contaminated equipment or materials.

Derra,⁹ recorded that the pregnant women are about 20 times more at risk than others and about 1/3 of listeriosis cases occur during pregnancy, late in 2^{nd} or in 3^{rd} trimester or 3 weeks of the newborn life . In pregnant women infection of the fetus is extremely common and can lead to abortion, still birth or delivery of Listeria infected infant. In Alberta, prevalence of L. monocytogenes was (30%) in pregnant women >40 years, and (22%) in non-pregnant women more than 40 years³⁵.

The current result were inconsistent with result found by Silk et al.,³² in Atlanta , whom they found that the crude prevalence of listeriosis among Hispanic pregnancy women were increased from 5.09 to 12.37 cases per 100 000 of population for the periods of 2004–2006 and 2007– 2009 respectively , while the prevalence of listeriosis among non-Hispanic pregnancy women were increased from 1.74 to 2.80 cases per 100.000 population for the same periods. Incidence rates of non-pregnancy associated listeriosis in patients aged \geq 65 years were 4–5 times greater than overall rates annually.

The results obtained in Table 10 and Figure 9 illustrated that the highest Prevalence rate of L. monocytogenes in total Human samples during period of study was in August , September and April (44.4%) , (34.3%) and (33.3%) respectively. The results indicated that there were significant differences at

(p< 0.05) in blood male and placenta, but there were no significant differences at (p> 0.05) in blood female.

In USA ³⁶ reported that the isolation of L. monocytogenes was higher during cooler weather (28 to 92% of samples) than during warmer weather (6 and 77% of samples). Also in Greece ³⁷ found that the prevalence of L. monocytogenes was higher in the warmer months. In Finland ³⁸ found that the seasonal trend seems evident in human listeriosis cases, and the number of these cases began to rise in July and remained high until January especially in August, September, October and January.

Data which has found clear seasonal variation, including higher numbers of Listeria species occurring in both winter and in summer months^{36, 39}. Also FDA, ⁴⁰ mentioned that infections with L. monocytogenes occur throughout the year. In Ireland³⁰ reported that the lowest prevalence occurred in July 2013, November 2013 and January 2014 (3.9, 3.8, and 2.0% respectively), while L. monocytogenes prevalence ranged between 4.2 and 6.0% for the rest of sampling months.

Depending on statistics collected from General Thamar Hospital (Table 11, Figure 10), we noticed that the recovered cases of abortion in this hospital during four years reach to 1072 cases, and the abortion cases recorded in all directorates, also this table clarify that the highest number of abortion cases were in Thamar Directorate compared with another directorates, our results agreement with the statistics achieved from the General Thamar Hospital, and this findings explain the risk of listeriosis.

CONCLUSION

Due to the importance of this study we recommend further researches and studies on L. monocytogenes in different governorates of Yemen, especially on the prevalence of L. monocytogenes in food of animal origin. Also working on creating specialized public centers deal with further studies and diagnosis of L. monocytogenes and emphasis for inter the diagnosis of L. monocytogenes within the routine work of the laboratories in hospitals. This study also recommends all people especially who do in field of food preservation and food cooking either in household or that working in restaurants to follow the following instructions: Keep clean, always wash hands with soap after going to the toilet. Separate raw and cooked food; avoid contacting between raw and cooked food and keep food at safe temperature. Cook thoroughly, especially meat and poultry. Use safe water and raw materials, wash fruit and vegetables.

REFERENCES

1- AHW (Alberta Health and Wellness). (2011): Listeriosis . Public Health Notifiable Disease Management Guidelines. Government of Alberta, Canada.

- 2- Ba-Salamah , H.A. (2013): Prevalence of Listeria monocytogenes in Human and Red meat and Study the Effect of Thermal Treatment on Isolated Bacteria in Thamar Province. M.SC. thesis, Faculty of Applied Sciences, Thamar University, Dhamar, Yemen.
- 3- Bula, C. J.; Bille, J. and Glauser, M. P. (1995): An epidemic of foodborne Listeriosis in western Switzerland: description of 57 cases involving adults . Clinical Infect Diseases Journal, 20 (1):66-72.
- 4- Cao, X.; Wang, Y.; Wang, Y. & Ye, C. (2017). Isolation and characterization of Listeria monocytogenes from the black-headed gull feces in Kunming, China. Journal of Infection and Public Health, 2017.
- 5- Cheesbrough, M. (2009): District Laboratory Practice in Tropical Countries. second Edition, part 2, Cambridge University Press, Cambridge, UK.
- 6- Cossart P. and Lebreton, A. (2014): A trip in the "New Microbiology" with the bacterial pathogen Listeria monocytogenes. FEBS Letters 1; 588(15):2437-45.
- 7- Derra, F. A. (2007): Prevalence and Antimicrobial Profile of Listeria monocytogenesin Retail Meat and Dairy Products in Addis Ababa and its Surrounding Towns, Ethiopia. Ph.D. Thesis, Department of Microbiology, Immunology, and Parasitology, Faculty of Medicine, Addis Ababa University, Ethiopia.
- 8- DOH (Washington State Department of Health): Listeriosis. Last Revised : January 2017, Page 1 of 7
- 9- ECDC (European Centre for Diseases Prevention and Control) (2017): Facts about listeriosis Stockholm: ECDC; 2017. Available online from: https://ecdc.europa.eu/en/listeriosis/facts.
- 10- Eslami ,G.; Samadi , R.; Taherpanah , R.; Taherpor , A.; Baseri , N. (2014) : Detection of actA and inlB genes in Listeria monocytogenes isolated from women with spontaneous abortions, Nov. Biomed. 2 (1) : 18e21.
- 11- FDA (U.S. Food and Drug Administration) (2011): Listeriosis (Listeria infection). multistate outbreak of Listeriosis-centers for disease control and prevention, Last Reviewed: October 2011.
- 12- Filipello, V.; Amato, E.; Gori, M.; Huedo, P.; Ciceri,G.; Lomonaco, S.; and Pontello,M. (2017): Epidemiology and Molecular Typing of Pregnancy-Associated Listeriosis Cases in Lombardy, Italy, over a 10-Year Period (2005–2014). Infectious Diseases in Obstetrics and Gynecology Volume 2017.
- 13- Filipello, V.; Amato, E.; Gori, M.; Huedo, P.; Ciceri , G.; Lomonaco , S. & Pontello, M. (2017): Epidemiology and Molecular Typing of Pregnancy-Associated Listeriosis Cases in Lombardy, Italy, over a 10-Year Period (2005–2014). Infectious diseases in obstetrics and gynecology, 2017.
- 14- Fugett, E. B.; Bopp, D. S.; Dumas, N. B.; Corby, J. and Wiedmann, M. (2007): Pulsed-Field Gel Electrophoresis (PFGE) Analysis of Temporally Matched Listeria monocytogenes Isolates from human clinical cases, foods, ruminant farms, and urban and natural environments reveals source-associated as well as widely distributed pfge types. Journal of Clinical Microbiology, Vol. 45, No. 3, p: 865–873.
- 15- Gillespie, I. A.;Mook, P.; Little, C. L.; Grant, K. A. and Mclauchlin, J. (2010): Human Listeriosis in England, 2001- 2007: association with neighborhood deprivation.

- 16- Gracey, J. F.; Collins, D. S. & Huey, R. J. (1999): Meat Hygiene. 10th Edition, W.B. Saunders Company Ltd. London, UK.
- 17- Guerini, M. N.; Harhay, D. M. B.; Shackelford, S. D.; Arthur, T. M.; Bosilevac, J. M.; Kalchayan, N.; Wheeler, T. L. and Koohmaraie, M. (2007): Listeria prevalence and Listeria monocytogens serovar diversity at cull cow and bull processing plants in the United States. Journal of Food Protection, 7 (88): 456 - 459.
- 18- Indrawattana,N.; Nibaddhasobon,T.; Sookrung , N.; Chongsa-nguan,M.; Tungtrongchitr , A.; Makino , S.; Tungyong , W. , and Chaicumpa , W. (2011) :Prevalence of Listeria monocytogenes in raw meats marketed in Bangkok and characterization of the isolates by phenotypic and molecular methods, J. Health Popul. Nutr. , 29 (1) : 26 38.
- Jensen, K.; Ethelberg, S.; Smith, B.; Nielsen, E. M.; Larsson, J.; Mølbak, K.; Christensen, J. J. and Kemp, M. (2010): Substantial increase in listeriosis. Rapid communications, Denmark.
- 20- Lake, R.; Hudson, A.; Cressey, P. and Gilbert, S. (2005): RiskProfile: Listeria monocytogenesin Soft Cheeses. Institute of Environmental Science & Research Limited, Christchurch Science Centre, New Zealand.
- 21- Larsen, C. N.; Nørrung, B.; Sommer, H. M. and Jakobsen, M. (2002): In Vitro and In Vivo Invasiveness of Different Pulsed-Field Gel Electrophoresis Types of Listeria monocytogenes. Applied and Environmental Microbiology, Vol. 68, No. 11, p: 5698–5703.
- 22- Leong, D.; Avelino Alvarez-Ordóñez, A. & Jordan , K. (2014): Monitoring occurrence and persistence of Listeria monocytogenes in foods and food processing environments in the Republic of Ireland .Front Microbiol.; 5: 436. Published online 2014 Aug 20.
- 23- Liu, D. (2008): Handbook of Listeria monocytogenes. CRC Press.
- 24- Lotfollahi, L.; Chaharbalesh, A.; Rezaee, M. A. & Hasani, A. (2017) : Prevalence, antimicrobial susceptibility and multiplex PCR-serotyping of Listeria monocytogenes isolated from humans, foods and livestock in Iran . Microbial Pathogenesis 107, 425-429
- 25- Lotfollahi, L.; Nowrouzi, J.; Irajian, G.; Masjedian, F.; Kazemi, B.; Eslamian, L.; Falahat, A. and Ramez, M. (2011): Prevalence and antimicrobial resistance profiles of Listeria monocytogenes in spontaneous abortions in humans. African Journal of Microbiology Research, Vol. 5 (14): 651-658.
- 26- Lukinmaa, S.; Miettinen, M.; Nakari, U. M.; Korkeala, H. & Siitonen, A. (2003): Listeria monocytogenes Isolates from Invasive Infections: Variation of Sero- and Genotypes during an 11-Year Period in Finland. Journal of Clinical Microbiology, Vol. 41, No. 4, p: 1694–1700.
- 27- Mammina, C.; Aleo, A.; Romani, C.; Pellissier, N.; Nicoletti, P.; Pecile, P.; Nastasi, A. & Pontello, M. M. (2009): Characterization of Listeria monocytogenesIsolates from human listeriosis cases in italy. Journal of Clinical Microbiology, Vol. 47, No. 9, p: 2925–2930.

- 28- Nazar, M. S. (2006): Study on Immuno pathological Changes Caused by Listeria monocytogenes in Mice and Lambs. M.Sc. Thesis, College of Veterinary Medicine, University of Baghdad, Baghdad, Iraq.
- 29- Nilsson, R. E. (2010): Studies on the biology of environmentally persistent Listeria monocytogenes strains. Ph.D. Thesis, University of Tasmania, Tasmania, Australia.
- 30- Rebagliati, V.; Philippi, R.; Rossi, M. and Troncoso, A. (2009): Prevention of foodborne listeriosis. Buenos Aires University, Argentina. Indian Journal of Pathology and Microbiology, Volume: 52, Page: 145-149.
- 31- Rhoades, J. R., G. Duffy, and K. Koutsoumanis. "Prevalence and concentration of verocytotoxigenicEscherichia coli, Salmonella enterica and Listeria monocytogenes in the beef production chain: a review." Food microbiology 26, no. 4 (2009): 357-376.
- 32- Rivoal , K.; Queguiner, S.; Boscher, E.; Bougeard , S.; Ermel , G.; Salvat G., et al. (2010): Detection of Listeria monocytogenes in raw and pasteurized liquid whole eggs and characterization by PFGE. Int. J. Food Microbiol. 138, 56–62
- 33- Rodrigues, C. S.; Cordeiro de Sá, C. V. G. and de Melo1, C. B. (2017): An overview of Listeria monocytogenes contamination in ready to eat meat, dairy and fishery foods. Ciência Rural, Santa Maria, v.47: 02, e20160721.
- 34- Saleh, M. G. S. (2010): Investigation on Salmonella spp. in patients suffering from fever and food poisoning in Thamar city- Yemen. M.Sc. Thesis, Sana'a University, Faculty of Science, Biology Department, Microbiology Section, Sana'a, Yemen.
- 35- Schuppler, M. and Loessner, M. J. (2010): The Opportunistic Pathogen Listeria monocytogenes: Pathogenicity and Interaction with the Mucosal Immune System. International Journal of Inflammation, Vol. 20. P: 1-12.
- 36- Silk, B. J.; Date, K. A.; Jackson, K. A.; Pouillot, R.; Holt, K. G.; Graves, L. M.; Ong, K. L.; Hurd, S.; Meyer, R.; Marcus, R.; Shiferaw, B. and Mahon, B. E. (2012): Invasive Listeriosis in the Foodborne Diseases Active Surveillance Network (Food net), 2004- 2009: Further Targeted Prevention Needed for Higher-Risk Groups. Clinical Infectious Diseases Journal, Vol. 54, No. 5, p: 396-404.
- 37- Sjoman, M. (2010): The use of Serotyping and PFGE-Typing of Listeria monocytogenesin food Processing Contamination Studies and Human Foodborne Infections. M.Sc. Thesis, Department of FoodHygiene and Environmental Health , Faculty of Veterinary Medicine, University of Helsinki, Finland.
- 38- Todd, E. C. D., &Notermans, S. (2011). Surveillance of listeriosis and its causative pathogen. Listeria monocytogenes. Food Control, 22, 1484e1490.
- 39- Weller , D.; Andrus , A.; Wiedmann , M. and den Bakker, H.C. (2015) : Listeria booriae sp. nov, and Listeria newyorkensis sp. nov., from food processing environments in the USA. International Journal of Systematic and Evolutionary Microbiology , 65 , 286 292 .

40- Zhu, Q; Gooneratne, R. & Hussain, M. A. (2017): Listeria monocytogenes in Fresh Produce : Outbreaks , Prevalence and Contamination Levels . Foods 2017, 6, 21.

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Test	Reaction
Catalase	+
Oxidase	-
Indole	-
Urease	-
Gram Stain	+
Motility 25°C	+
37°C	-
H ₂ S production	-
Hemolytic (β)	+
TSI	A/A
Methyl red	+
Simmon's citrate	-
Voges- proskaur	+

Table (1): Biochemical Tests for the Confirming of L. monocytogenes

Table (2): Prevalence of L. monocytogenesin Total Human Samples

Samples	Total N. of Samples	Positive samples %No	Negative samples %No	p-value
Male Blood	100	18 18.0	82 82.0	
Female Blood	110	22 20.0	88 80.0	0.862
Placenta	100	21 21.0	79 79.0	
Total	310	61 19.7	249 80.3	



Figure (1): Prevalence of Listeria monocytogenes in Total HumanSamples

Say	Total No. of	Posi	tive	Neg	ative
Sex	Samples	No.	%	No.	%
Male Blood	100	18	18.0	82	82.0
Female Blood	110	22	20.0	88	80.0
Total	210	40	19.0	170	80.9





Figure (2): Prevalence of *L. monocytogenes* in Human according to Sex

Table (4): Prevalence of L.monocytogenesin male according to Directorates Prevalence

Directorates	Total N. of	Pos	itive	Ne	egative	p-value
Directorates	Samples	N.	%	N.	%	
Thamar	35	35 11 31.4 24 68.6				
Alhada	22	2	9.1	20	90.9	
Ans 28		3	10.7	25	89.3	
Anes	2	0	0.0	2	100.0	.138
Gahran	13	2	15.4	11	84.6	
Total	100	18	18.0	82	82.0	





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	Total N. of	al N. of Positive			gative	p-value
Directorates	Samples	N.	%	N.	%	ļ •
Thamar	34	10	29.4	24	70.6	
Alhada	28	5	17.9	23	82.1	
Ans	24	3	12.5	21	87.5	E 4 2
Anes	10	2	20.0	8	80.0	.545
Gahran	14	2	14.3	12	85.7	
Total	110	22	20.0	88	80.0	

Table (5): Prevalence of *L.monocytogenes*in female according to Directorates



Figure (4): Prevalence of *L.monocytogenes*in Female according to Directorates

	Total N. of	tal N. of Positive		Ne	gative	p-value
Directorates	Samples	N.	%	N.	%	
Thamar	37	2	5.4	35	94.6	
Alhada	27	9	33.3	18	66.7	
Ans	19	5	26.3	14	73.7	004
Anes	6	1	16.7	5	83.3	.004
Gahran	11	4	36.4	7	63.6	
Total	100	21	21.0	79	79.0	

Table (6): Prevalence of *L.monocytogenes*in Placenta according to Directorates

Prevalence of Listeria monocytogenes in Human in Dhamar Governorate/ Yemen

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Figure (5): Prevalence of L.monocytogenesin Placenta according to Directorates

Distribution of samples by	N of Samples	Ро	sitive	Neg	gative	
Age	IN. OF Samples	N.	%	N.	%	p-value
< 20	27	3	11.1	24	88.9	
21-30 years	years 24		12.5	21	87.5	
31-40 years	18	3	16.7	15	83.3	0 537
41-50 years	11	3	27.3	8	72.7	0.337
≥60	≥ 60 20		30.0	14	70.0	
Total	100	18	18.0	82	82.0	

Table	(7): Prevalence	of Listeria mono	cvtogenesin Ma	le Blood acco	ording to Age
	(.)				



Figure (6): Prevalence of Listeria monocytogenesin Male Blood according to Age

		, ,			6	0 0		
Distribution of Samples	N of Samples	Posit	ive	Neg	gative	n-value		
by Age		N.	%	N.	%	p ⁻ value		
Less than 20	26	2	7.7	24	92.3			
21-30 years	44	12	27.3	32	72.7			
31-40 years	19	3	15.8	16	84.2	494		
41-50 years	12	3	25.0	9	75.0	.404		
≥51	9	2	22.2	7	77.8			
Total	110	22	20.0	88	80.0			

Table (8): Prevalence of *Listeria monocytogenes* in female blood according to ages



Figure (7): Prevalence of *Listeria monocytogenes* in female blood according to ages

Table (9): Prevalence of Listeria monocytogenespregnancy woman accordingto age in Thamar province

Distribution of Samulas by Aga	N. of	Pos	itive	Neg	gative	n valua
Distribution of Samples by Age	Samples	N.	%	N.	%	p- value
Less than 20	17	5	29.4	12	70.6	
21-30 years	48	7	14.6	41	85.4	
31-40 years	33	9	27.3	24	72.7	.357
41-50 years	2	0	0.0	2	100.0	
Total	100	21	21.0	79	79.0	

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 Table (10): Relationship between months and prevalence of L.monocytogenesintotal human

 samples during period of study

м		ths	No, of Male Blood Femal		e Blood	Plac	enta	Total of Pos. Samples			
			Samples	N.	Pos.	N.	Pos.	N.	Pos.	N.	%
	Febru	ıary	35	12	0	11	1	12	1	2	5.7
	Mai	·ch	35	12	0	12	1	11	0	1	2.9
April		ril	36	12	6	12	1	12	5	12	33.3
May		34	11	1	13	4	10	3	8	23.5	
June			36	11	0	13	2	12	3	5	13.9
	Jul	у	34	11	1	13	2	10	1	4	11.8
	Aug	ust	36	10	7	13	7	13	2	16	44.4
	Septer	mber	35	11	3	11	3	13	6	12	34.3
	October 29 10 0 12 1				7	0	1	3.5			
	Tot	al	310 100 18 110 22 10		100	21	61	19.7			
p- value	0.001	.0620	0.004								





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Table (11): Relationship between Directorates and abortion cases in General Thamar Hospital from2009 - 2012.

Years Months	2009	2010	2011	2012	Total
January	0	8	22	22	52
February	0	10	35	26	71
March	0	17	43	26	86
April	19	13	31	30	93
May	12	9	38	37	96
June	5	14	20	34	73
July	15	24	16	37	92
August	4	20	27	32	83
September	3	14	30	35	82
October	10	8	34	56	108
November	0	20	30	63	113
December	13	43	29	38	123
Total	81	200	355	436	1072





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مدى انتشار بكتيريا Listeria monocytogenes في الإنسان في محافظة ذمار/ اليمن وذلك تبعاً للفئات العمرية المختلفة

الملخص: أجربت هذه الدراسة لتحديد مدى انتشار بكتيريا Listeria monocytogenes في الإنسان في محافظة ذمار/ اليمن وذلك تبعاً للفئات العمرية المختلفة.

تم جمع 310 عينة من الإنسان وبواقع 100 عينة دم من الذكور، 110 عينة دم من الإناث، 100 عينة من المشيمة، حيث جمعت العينات من مستشفى ذمار العام ومستوصف المصلي والمختبرات الطبية في المحافظة .

أظهرت نتائج الدراسة بأن النسبة الكلية لانتشار بكتريا L. monocytogenes من إجمالي العينات كانت61 (19.7%)، وأن نسبة عزل البكتريا من دم الإناث 22 (% 20) كانت أعلى من دم الذكور 18(%18.)، وأما من المشيمة فقد بلغت 21 (21%).

كما أشارت النتائج التي تخص نسبة عزل الجرثومة من دم الإنسان حسب المديريات إلى أن نسبة العزل في الذكور كانت 11 (4.0%) في مديرية ذمار، 2 (15.4%) في مديرية قاس، 2 (1.9%) في مديرية الحداء، بينما لم تعزل أي عينة من مديرية عنس. كما يتضح لنا بان نسبة العزل في الإناث كانت 10 (4.9%) في مديرية ذمار، 2 (2.0%) في مديرية عنس، 5 عينة من مديرية عنس. كما يتضح لنا بان نسبة العزل في الإناث كانت 10 (4.9%) في مديرية ذمار، 2 (0.0%) في مديرية عنس، 5 (1.0%) في مديرية قابرات 10 (4.9%) في مديرية الحداء، بينما لم تعزل أي عينة من مديرية عنس. كما يتضح لنا بان نسبة العزل في الإناث كانت 10 (4.9%) في مديرية ذمار، 2 (0.0%) في مديرية عنس، 5 العزل في الإناث كانت 10 (4.9%) في مديرية ذمار، 2 (0.0%) في مديرية عنس، 5 (1.9%) في مديرية عنس، 5 (1.9%) في مديرية عنس، 5 (1.9%) في مديرية محبران، 3 (2.5%) في مديرية انس. اما نسبة عزل الجرثومة من المشيمة حسب المديريات، فقد أشارت النتائج إلى أن أعلى نسبة لعزل الجرثومة كانت في مديرية جهران 4 (4.5%)، وتلتها مديرية الحداء، 9 (3.5%) في مديرية جهران، 3 مديرية جهران، 3 مديرية الحداء، 2 (3.5%) في مديرية عنس. المديريات، فقد أشارت النتائج إلى أن أعلى نسبة لعزل الجرثومة كانت في مديرية جهران 4 (4.5%)، وتلتها مديرية الحداء 9 (3.5%)، مديرية آنس(3.6%)، وتلتها مديرية الحداء 9 (3.5%)، مديرية آنس(3.6%)، مديرية منسرة 1.5%) ومديرية ذمار (4.5%). وأظهرت الدارسات الاحصائية فروق ذات دلالة معنوية عند مديرية آنس(3.6%)، مديرية منسرة 1.5%)، ومديرية دمار (4.5%). وأظهرت الدارسات الاحصائية فروق ذات دلالة معنوية عند (p<0.05) بين انتشار p<0.05%) بين انتشار p<0.05%) ومديرية دمار حسب المديريات.

وعند دراسة العلاقة بين المراحل العمرية المختلفة وبين نسبة عزل البكتريا، وجدنا أن أعلى نسبة للعزل من دم الذكور كانت في الفئة العمرية التي تزيد عن (60) سنة بنسبة (30,0%)، وتلتها الفئة العمرية التي تتراوح ما بين (41- 50) سنة بنسبة (2.72%)، ثم الفئة العمرية (31-40) سنة بنسبة (16.7%)، والفئة العمرية (21-30) سنة بنسبة (2.51%)، وأخيراً الفئة العمرية التي تقل أعمارها عن(20) سنة بنسبة (11.1%). بينما كانت أعلى نسبة عزل من دم الإناث حسب الفئات العمرية تراوحت ما بين (20) سنة بنسبة (2.72%)، ثم تلتها الفئات العمرية (11.1%). بينما كانت أعلى نسبة عزل من دم الإناث حسب الفئات العمرية تراوحت ما بين (21-30) سنة بنسبة (2.72%)، ثم تلتها الفئات العمرية (2.20%)، والفئة العمرية التي تزيد عن(51) سنة و(30-40) سنة وأخيراً الفئة العمرية التي تقل عن(20) ثم تلتها الفئات العمرية (21.0%)، والفئة العمرية التي تزيد عن(51) سنة و(30-40) سنة وأخيراً الفئة العمرية التي تقل منة بنسبة (2.0%)، (2.22%)، (8.75%) و (7.7%) على التوالي. وقد أوضحت النتائج أن أعلى نسبة إصابة في النساء الحوامل كانت في الفئة العمرية التي تقل عن (20) سنة بنسبة (2.9%)، وتلتها الفئة العمرية العمرية التي توام كانت في منة بنسبة (2.5%)، وعزلة من الفئة العمرية القي تزيد عن(51) سنة و(3.40) سنة وأخيراً الفئة العمرية التي تقل عن (2.6%)، وعلي تقل عن (20) منة بنسبة (2.7%)، وتلتها الفئة العمرية العمرية التي وقد أوضحت النتائج أن أعلى نسبة إصابة في النساء الحوامل كانت في (1.6%)، ولم تعزل أي عزلة من الفئة العمرية ما بين (4.0%)، سنة الفئة العمرية (3.40%)، شم (12-30%)، ما قد وبنسبة

أما النتائج التي تخص العلاقة ما بين نسبة انتشار البكتريا في العينات المأخوذة من الإنسان وما بين أشهر السنة ، فقد أظهرت أن أعلى نسبة كانت في شهر اغسطس وبنسبة(44.4%) وشهر سبتمبر وبنسبة (34.3%) .

استنتج من هذه الدراسة أن نسبة انتشار بكتريا L. monocytogenes في محافظة ذمار كانت عالية، وأن إصابة الإنسان بهذه الجرثومة يمكن أن تحدث في أية مرحلة من مراحل العمر وبخاصة المراحل المتقدمة من العمر في الذكور والمراحل التي تزداد فيها نسبة الحمل في الإناث.

أوصت الدراسة بضرورة تواصل الأبحاث والدراسات عن بكتيريا L. monocytogenes في مختلف محافظات الجمهورية اليمنية، والعمل على إنشاء مراكز وطنية متخصصة في مجال عزل وتشخيص هذه البكتيريا. كما أكدت على أهمية نشر التوعية الصحية عن طريق وسائل الاعلام المرئية والمسموعة والمقروءة ولكافة شرائح المجتمع من الذكور والإناث، وتسليط الضوء على المخاطر الصحية الناتجة عن الإصابة بهذه البكتيريا، واتباع السياقات العلمية المعتمدة في مجال الصحة العامة لمنع انتقالها إلى الإنسان.

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Prevalence of Listeria monocytogenes in Human in Dhamar Governorate/ Yemen