# THE INTERNATIONAL JOURNAL OF SCIENCE & TECHNOLEDGE

# **Microbial Quality of Carcass in Khartoum State**

### Hamid El nourain Hamdan

Assisstant Professor, Department of Food Safety, Faculty of Public Health, AlZaiem Alazhari University, Sudan

#### Abstract:

A total of 60 random swab samples were collected from cattle, camel and sheep carcasses at Omdurman slaughter houses to evaluate the contamination level with Enterobacteriaceae. The results obtained indicated that the mean values of bacterial counts in the examined swab samples of cattle, sheep and camel were 3.6, 3.13 and 2.72 Mean Log<sub>10</sub> cfu/cm<sup>2</sup> and 3.4,2.9 and 2.34 Mean Log<sub>10</sub> cfu/cm<sup>2</sup> for the total coliform count, respectively. There was a significant difference between Enterobacteriaceae and coliform counts. E.coli (20%), Citrobacter (10%), Enterobacter (35%), Klebsiella (25%) and Proteus species(30%) ,were the predominant bacterial species among the isolates. From all the three types of carcasses No salmonella species were detected in any sample.

Keywords: Carcass, slaughter- house, coli forms

#### 1. Introduction

Fresh meat is highly perishable due to its biological components. Microbial contamination of the carcass during the slaughtering process results in spoilage of meat, reduced shelf-life of meat and public health hazards (Narasimha Rao, 1992), (Nortje, Nel, Jordaan, badenhorst, Goedhart, & Holzapfel, 1990). Almost most food borne diseases are related to use of meat containing pathogenic microorganisms. There is high possibility of external contamination of carcasses i from the early stages of slaughtering till the moment consumption. Good manufacturing practices during slaughter have a profound effect on the microbial load of meat. Contact with the hide, skin or feet, and the gastric content are considered as potential sources for meat contamination by microorganisms. Water sources, air, instrument used for slaughter operations such as (knives, saws, cleavers or hooks) play a role in meat contamination(Jawelz, E.J.; Meluick, L.and E.A.Adelbery., 1982). The Enterobacteriaceae family contains a large number of organisms, some of non faecal origin, that are useful as an indicator of the overall process hygiene in the abattoir. E. coli is the indicator bacteria of choice associated with faeces (Delhalle, de Sadeleer, Bollaerts, Farnir, & Saegerman, 1994) (Ghafir, China, Dierick, & De Zutter, 2008) and (McEvoy, Sheridan, & Blair, 2004) Enteric organisms, such as coliforms were frequently isolated from meat indicating that the rumen of the slaughtered animals is a common source of contamination (ICMSF I. C., 1980). Therefore, the objective of the current study was to determine the level of Enterobacteriaceae contamination of sheep, cattle and camel carcasses during slaughtering and to identify their pathogenic strains.

# 2. Materials and Methods

# 2.1. Collection of Samples

A total of 60 random swab samples were collected from slaughtered cattle, camel and sheep carcasses at Omdurman slaughter houses, using a sterile wire template with an approximate area of 100 cm<sup>2</sup>. The swabs were delivered immediately to the laboratory for testing. Samples were collected according to(ICMSF I. C., 1978)

### 2.2. Bacterial Counts

Enterobacteriaceae count was determined as by using Violet Red Bile Glucose Agar and the purple colonies were calculated as Enterobacteriacae. For the Determination of coliform count Violet Red Bile agar plates were used (ICMSF.International Commission on Microbiological Specification for Foods., 1996).MacConkey broth and Eosin Methylene Blue plates were used for detection of *E.coli*. The metallic green colonies were picked up and identified biochemically (ICMSF.International Commission on Microbiological Specification for Foods., 1996). For salmonellae detection peptone water was used as pre enrichment medium, Rappaport Vassiliadis as an enrichment broth and Hekton agar was used for plating.

#### 2.3. Statistical Analysis

Bacterial counts were transformed into log cfu/cm<sup>2</sup>values.data were tested using SPSS version 16(SPSS inc.Chicago). Turkeys' test was used as a post hoc test. Mean differences were considered significant at p< 0.05. Data was represented in form of tables and graphs.

#### 3. Results

The International Journal Of Science & Technoledge

Carcasses	Positive samples		Mean Log <sub>10</sub> cfu/cm <sup>2</sup>
	No	%	
Cattle	16	80	$3.6^*$
Sheep	8	40	3.13
Camel	5	25	2.72

Table 1: Mean  $Log_{10}$  cfu/cm<sup>2</sup> of Total Enterobacteriaceae counts recovered from different animal carcasses (n=20). \*Significant difference ( $P \le 0.05$ )

Carcasses	Positive	samples	Mean Log <sub>10</sub> cfu/cm <sup>2</sup>
	No	%	
Cattle	18	90	3.4**
Sheep	15	75	2.90
Camel	12	60	2.34

Table 2: Mean  $Log_{10}$  cfu/cm<sup>2</sup> of Total coliform counts recovered from different animal carcasses (n=20). \*\* High Significant difference  $(P \le 0.01)$ 

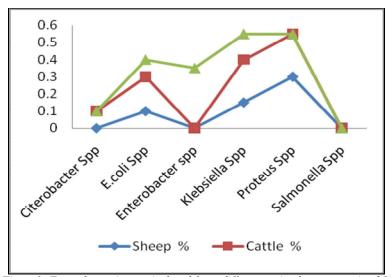


Figure 1: Enterobacteriaceae isolated from different animal carcasses (n=20)

#### 4. Discussion

The obtained results in table (1) and (2), indicate that the total Enterobacteriaceae count in the examined swab samples were varied from 2.72 to 3.4 Log<sub>10</sub> cfu/cm<sup>2</sup> with an average of 3.4 Log<sub>10</sub> cfu/cm<sup>2</sup> for cattle, 3.13 Log<sub>10</sub> cfu/cm<sup>2</sup> for sheep and 2.72 Log<sub>10</sub> cfu/cm<sup>2</sup> for camel. Significant differences were detected among different species of carcasses at  $(P \le 0.05)$ . Nearly similar results were obtained by (Hamdy, 1989), (Samaha, 1993). Higher results were obtained by (Al-dughaym, 2001), concerning the mean values of Enterobacteriaceae count on the surface of camel carcasses before and after skinning, and at the inspection point. While, lower results were obtained by (Pearce, 2005) and (Fliss & Simrad, 1991), (Vanderlinde, 1999), (Yalcin & Nizamlioglu, 2001) during the different stages of slaughter operations. Regardless of animal species; counts were relatively higher for freshly produced meat.

Different species of microorganisms were isolated from the samples, as E,coli & Klbsiella were isolated from all animal species carcasses., Proteus from sheep and cattle, while Citrobacter was isolated from cattle carcasses only, all species carcasses were found free of Salmonella Figure (1). All examined swabs from the three species carcasses were free from Salmonellae. This finding agrees with some researchers, who could not detect Salmonellae spp., from any of the examined samples, although (WHO, 1988) records showed that, the incidence of Salmonella in raw meat and organs in some countries was up to 10% and 3%, respectively, globally Salmonella is implicated in most food poisoning outbreaks occurring as a result of consumption of contaminated meat and meat products. Members of the Enterobacteriaceae are responsible for causing foodborne disease and some also cause food spoilage and therefore contribute to substantial economical losses and food wastage. The initial Enterobacteriaceae contamination level in the raw materials is predominantly governed by Good Agricultural Practices (GAP) during primary production and subsequently during slaughter of livestock at the abattoir (Chris Baylis, 2011).

# 5. Acknowledgements

I would like to thank the administration and workers of the slaughter houses at Khartoum State for their cooperation, and my thanks are also due to the Environmental Health authorities for their appreciated help.

#### 6. References

- 1. Abou-Yossef, H. M. (2010). Quality assurance of camels meat. Alexandria: Faculty of Veterinanry Science, Alex, Univ.
- 2. Al-dughaym, A. a. (2001). Surface contamination of camel carcasses. Basic and Applied Sciences. 2(1): , 129-138.
- 3. Chris Baylis, M. U. (2011). The Enterobacteriacae and Thier Significance To Food Industry. Brussels: ILSI.
- 4. Delhalle, L., de Sadeleer, L., Bollaerts, K., Farnir, F., & Saegerman, C. k. (1994). Risk factors for salmonella and hygiene indicators in the 10 largest Belgian pig slaughterhouses. Journal of Food Protection, 1320-1329.
- 5. Fliss, I., & Simrad, R. e. (1991). Microbial Quality of different fresh meats species in Tunisian slaughter houses and markets. J.Food Prot.54(10): ,773-777.
- 6. Ghafir, Y., China, B., Dierick, K., & De Zutter, L. D. (2008). Hygiene indicator microorganisms for elected pathogens on beef pork and poultry meats in Belgium. Journal of Food Protection 71, 35-45.
- 7. Hamdy, M. (1989). Surface contaminants of slaughtered camels. Zag.Vet.J.17(3), , 291-302.
- 8. ICMSF, I. C. (1980). Microbial Ecology of foods. Volume 1. factors affecting life and death of microorganisms. Orlando: Academic Press.p.311.
- 9. ICMSF, I. C. (1978). Microrganisms in Foods, Their significance and enumeration. 2nd ed. Toronto and Buffalo, Canada: Un. of Toronto Press.
- 10. ICMSF.International Commission on Microbiological Specification for Foods. (1996). Microrganisms in Foods.volume 5.Characteristics of microbial pathogens. London: Blackie Academic&Professional.
- 11. Jawelz, E., & Meluick, L. E. (1982). Review of Medical Microbiology, 16th Ed. Middle east Edition.
- 12. Jawelz, E.J.; Meluick, L.and E.A.Adelbery. (1982). Review of Medical microbiology.16th Ed.
- leung, P., Yam, W., & Ng, W. P. (2001). The and pigs in prevalence and characterization of verotoxin-producing Echerichia coli isolated form cattle and pigs in an abattoir in Hnog Kong. from cattle. Epidemiology and Jnfection, 126, , 173-179.
- 14. McEvoy, J., Sheridan, J., & Blair, I. M. (2004). Microbial contamination on beef in relation to hygiene assessment based on criteria used in EU Decision 2001/471/EC. International Journal of Food microbiology, 92, 217-225.
- 15. Narasimha Rao, D. R. (1992). The microbiology of sheep carcasses processed in a modern Indian abattoir. Meat Sci,32, , 425-436.
- 16. Nortje, G., Nel, L., Jordaan, E., badenhorst, k., Goedhart, G., & Holzapfel, W. G. (1990). Aquantitative survey of a meat production chain tto determine the microbial profile of the final product. J,Food prot.53, , 411-417.
- 17. Pearce, R. B. (2005). Excision vs sponge swabing a comparison of methods for the microbiological sampling of beef,pork and lamb carcasses. Jornal of Applied Microbiology, v. 98, , 896-900.
- 18. Samaha, I. D. (1993). Air and water as sources of bacterial contamination of beef carcasses. Alex.J. Vet. Sci. 9(2), , 83-88.
- 19. Terrance, M., Barkocy-GALLAGHER, G., & Rivera-Betancourt, M. K. (2002). Prevalence and charchterization of non-O157 shiga toxin producing Escherichia coli on carcassesin commercial beef cattle processing plants,. Appl.Environ.microbiol.68, , 4847=4852.
- 20. Vanderlinde, P. (1999). Microbiological status of Australian sheep meat. Journal of food Protection, Ames, v. 2(4), , 380-385.
- 21. WHO, W. H. (1988). Salmonellosis control .The role of animal and product hygiene. Barking, Essex, UK.: Applied Sci. publishers, Ripple Road.
- 22. Woerner, D., Ransom, J., Sofos, J., Dewell, G., Smith, G., & Salman, M. a. (2006). Determining the prevalence of Escherichia coli O157 in cattle and beef from the feedlot to the cooler. Journal of Food Protection, 69(12), , 2824-2827.
- 23. Yalcin, S., & Nizamlioglu, M. a. (2001). Fecal coliform contamination of beef carcasses during the slaughtering process. Journal of Food safety, 21, , 225-231.