# Evaluation of Bacterial Composition of the Aerosol of Selected Abattoirs in Akure, South Western Nigeria

Funmilola O. Omoya, Joseph O. Obameso, Titus A. Olukibiti

Abstract-This study was carried out to reveal the bacterial composition of aerosol in the studied abattoirs. Bacteria isolated were characterized according to microbiological standards. Factors such as temperature and distance were considered as variable in this study. The isolation was carried out at different temperatures such as 27°C, 31°C and 29°C and at various distances of 100meters and 200meters away from the slaughter sites. Result obtained showed that strains of Staphylococcus aureus, Escherichia coli, Bacillus subtilis. Lactobacillus alimentarius and Micrococcus sp. were identified. The total viable counts showed that more microorganisms were present in the morning while the least viable count of 388cfu was recorded in the evening period of this study. This study also showed that more microbial loads were recorded the further the distance is to the slaughter site. Conclusively, the array of bacteria isolated suggests that abattoir sites may be a potential source of pathogenic organisms to commuters if located within residential environment.

Keywords—Abattoir, Aerosol, Bacterial Composition, Environment.

# I. INTRODUCTION

A n abattoir otherwise known as slaughterhouse is a facility where animals are killed for consumption as food products. It can be defined as livestock producing industries comprising operations relating to animals processing [11]. The major activities involved in the operation of an abattoir according [4] carcass boning and packaging, drying of skin and transport of processed materials.

Odor (air borne waste) is a major problem associated with abattoirs. Operations such as cooking and rendering process (include stale materials and fugitive emissions from cookers), waste effluent treatment plant, waste disposal technique such as burning dead stock animal holding pens (produced by manure and urine), odors from blood, skin handling and skin shed have been potential sources of pollution. Materials burned at an abattoir include coal or gas for boilers and steam production, diseased animals, sludge etc. Also, fuel burning which gives rise to atmospheric emissions is a source of pollution.

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Bio-aerosols are usually defined as aerosols or particulate matter of microbial, plant or animal origin that is often used as synonymously with organic dust. Bio-aerosols or organic dust may consist of pathogenic or non-pathogenic live or dead bacteria, fungi, viruses, high molecular weight allergens, plant fibers, mycotoxins, peptidoglycans,  $\beta$  (1 $\rightarrow$ 3)-glucansete [3].

The interest in bio-aerosol exposure has increased over the decades. This is largely because it is now appropriately recognized that exposure to biological agents in both residential and occupational indoor environment are associated with a wide range of adverse health effects with major public health impact. The diseases associated with bio-aerosol can be grouped into infectious diseases, respiratory diseases and cancer. Other adverse health effects as reported by [2], [7] include dermatitis in latex workers and pre-term birth or late abortion in farm women exposed to mycotoxins with immunetoxic and hormone-like effects. In this study, the bacterial composition of aerosol in and around abattoir sites is however investigated to assess the potential health effect of such facility in the environment.

### II. METHODOLOGY

#### A. Collection of Samples

Two major abattoir sites were visited trice for the collection of samples. They include Onyearugbulem slaughter house, Akure and the slaughter house at The Federal University of Technology, Akure Junction. The samples were collected aseptically by exposing agar plates of both general and selected culturing media such as nutrient agar, Eosin Methylene blue agar, and Mannitol salt agar to the abattoir air at the slaughter line and some meters away (100meters and 200meters) from the abattoir sites. These plates were exposed both in the morning  $(27^{0}C)$ , afternoon  $(31^{0}C)$  and evening  $(29^{0}C)$  thereafter were transferred into the laboratory and incubated at  $37^{0}C$  for 24 hours. Meanwhile the plates were coded with identity.

## B. Isolation of Microorganisms from Aerosols

The resultant colonies at the end of incubation were counted to know the total viable count and sub cultured to obtain pure cultures. Gram staining was carried out on the subculture to ascertain purity, which were thereafter transferred into double strength medium slants for further studies and identification. Cultural characterization of colonies; color, edge, elevation,

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surface and physiological and biochemical characterization tests using conventional methods [6] were employed.

### III. RESULTS AND DISCUSSION

The total viable count relatively varied with the different locations at the abattoir site (Table I). The abattoir B has the highest CFU at the slaughter line (606 cfu) as compared to Abattoir A (520 cfu). However, a relatively high value was recorded at some meters away from the abattoir site. At B<sub>L2</sub> (200 meters from the abattoir site) more bacterial count of 685 cfu was recorded. The closer the distance to the abattoir the lesser the total viable count obtained. The effect of temperature on the total viable count was considered in this study. At different temperature range, the total viable count differs (Table I). At low temperature (morning (27°C) and evening (29°C)) number of isolated microorganisms is a bit high compared to when the temperature is high. This could be as a result of high rate of degradation which reduces the bacterial load of some specific microorganisms which might not be able to adapt to the sudden change of environmental condition.

The presence of some of the isolated microorganisms in the abattoir is hazardous to human. The tests carried out showed biodiversity of both Gram positive and Gram negative pathogenic and non-pathogenic bacteria in the abattoir sampled (Table II). The species of bacteria were identified as Staphylococcus aureus, Escherichia coli, Bacillus subtilis, Lactobacillus alimentarius and Micrococcus sp. Lactobacillus sp. are known to cause spoilage in meat. Lactic acid bacteria can grow and spoil the meat when the oxygen tension in the package is low [8]. Lactobacillus alimentarius is reported to be a potential airborne meat spoilage microorganism and should be considered a processing critical control point [9], [5]. Escherichia coli is a bacterium responsible for outbreak of diarrhea. It is common in gastro intestine of warm blooded animals. Their presence is attributed to poor hygiene. This bacterium is known to cause food poison when taken in and result into gastro intestinal disease [12]. Staphylococcus aureus can cause a wide variety of diseases in human and other animals through toxin production. Staphylococcal toxins are a common cause of food poisoning, as they can be

produced by bacteria growing in improperly stored food items [1]. *Bacillus subtilis* is also known to cause disease in severely immunocompromised patients. It causes food poisoning and chronic skin infection [10]. In conclusion, the potential health effects of bio-aerosol from abattoir can be overwhelming; hence, the establishment of abattoir in residential environment can be a potential threat to humanity.

ACTERIAL	COUNT	(CFU)	) OF SAMPLED	ABATTO	IR SITES	AT DIFFE	ERENT PE	RIOD OF 7	THE DAY
				-		( 0)			

Samula	Total Viable Count(cfu)						
Sample	Morning (27°C)	Afternoon (31°C)	Evening (29°C)				
$A_{SL}$	520	450	388				
$A_{L1}$	600	560	520				
$A_{L2}$	633	592	542				
$\mathbf{B}_{\mathrm{SL}}$	606	557	475				
$B_{L1}$	674	565	545				
$B_{L2}$	685	624	566				

Legends

ASL- abattoir at the junction of federal university of technology, Akureslaughter line

AL1- abattoir at the junction of federal university of technology, Akure100meters from slaughter line

AL2- abattoir at the junction of federal university of technology, Akure200meters away from slaughter line

BSL- abattoir at Onyearugbulemmarkrt, Akureslaughter line

B

BL2- abattoir at the Onyearugbulemmarket, Akure100 meters away from abattoir site

BL2- abattoir at the Onyearugbulemmarket, Akure200 meters away from abattoir site

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Sample Locations	Isolates						
	S. aureus	L. ailmentarius	B. subtilis	E. coli	Micrococcus sp		
A <sub>MSL</sub>	+	+	-	+	+		
$A_{ML1}$	+	-	+	+	+		
$A_{ML2}$	+	+	+	+	-		
A <sub>NSL</sub>	+	+	+	+	-		
$A_{NL1}$	+	+	+	-	+		
A <sub>NL2</sub>	+	+	+	-	-		
$A_{ESL}$	+	-	+	+	-		
$A_{EL1}$	+	+	+	+	+		
$A_{EL2}$	+	+	+	+	-		
$B_{MSL}$	+	-	+	+	-		
$B_{ML1}$	+	-	+	+	+		
$B_{ML2}$	+	+	+	+	-		
B <sub>NSL</sub>	+	+	+	+	-		
B <sub>NL1</sub>	+	+	+	+	-		
B <sub>NL2</sub>	+	-	+	+	+		
$B_{ESL}$	+	-	+	+	+		
B <sub>EL1</sub>	+	+	+	+	-		
Brita	+	+	+	+	_		

 TABLE II

 FREQUENCY OF OCCURRENCE OF BACTERIA ISOLATES AT DIFFERENT PERIODS AND LOCATIONS

AMSL-morning samples from abattoir at the junction of Federal University of Technology, Akure slaughter line, AM L1- morning samples from abattoir at the junction of Federal University of Technology, Akure 200 m away from abattoir site, ANSL - afternoon samples from abattoir at the junction of Federal University of Technology, Akure 200 m away from abattoir at the junction of Federal University of Technology, Akure 200 m away from abattoir at the junction of Federal University of Technology, Akure 100 M Away From Abattoir at the junction of Federal University of Technology, Akure 100 M Away From Abattoir at the junction of Federal University of Technology, Akure 200 m away from abattoir at the junction of Federal University of Technology, Akure 200 m away from abattoir at the junction of Federal University of Technology, Akure 200 m away from abattoir at the junction of Federal University of Technology, Akure 100 m Away From Abattoir at the junction of Federal University of Technology, Akure 100 m away from abattoir at the junction of Federal University of Technology, Akure 200 m away from abattoir at the junction of Federal University of Technology, Akure 200 m away from abattoir at the junction of Federal University of Technology, Akure 100 m away from abattoir at Onyearugbulem market, Akure 100 m away from abattoir at Onyearugbulem market, Akure 100 meters away from abattoir at Onyearugbulem market, Akure slaughter line, BML1- morning sample from abattoir at Onyearugbulem market, Akure slaughter line, BNL1- afternoon sample from abattoir at Onyearugbulem market, Akure 100 meters away from abattoir at Onyearugbulem market, Akure slaughter line, BNL1- afternoon sample from abattoir at Onyearugbulem market, Akure 200 meters away from abattoir at Onyearugbulem market, Akure slaughter line, BNL1- afternoon sample from abattoir at Onyearugbulem market, Akure 100 meters away from abattoir at Onyearugbulem market, Akure slaughter line, BEL2-evening sample from abattoir at Onyearugbulem market, Akure 100 meters away

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#### REFERENCES

- [1] Centers for Disease Control and Prevention, 2010
- [2] Charous, B. L., Hamilton, R. G. and Yunginger, J. W. (1994). Occupational latex exposure: characteristics of contact and systemic reactions in 47 workers. *Journal of Allergy and Clinical Immunology*. 94: 12-8. Crawford.
- [3] Douwes, J., Thorne, P., Pearce, N and Heederik, D. (2003) Oxford Journals. Life Sciences and Medicine; *the annals of Occupational Hygiene*. 47: (3) 187-200
- [4] Grandin, T and Deesing, M (2008). Humane Livestock Handling. Storey publishing, North Adams, Ma, USA. Pp 67
- [5] Holley RA, MD Peirson, J Lam, KB Tan. (2004). Microbial profiles of commercial, vacuum packaged, of fresh pork of normal or short storage life. *Int J Food Microbiology*.
- [6] Holt, J. G., Krieg, N. R., Sneath, P. H. Stanley, J.J and Williams S.T. (1994) Bergey's Mannual of determinative Bacteriology. Wilkins Publishers. Baltimore.
- [7] Kristensen, P., Andersen, A. and rgens, L. M. (2000) Hormonedependent cancer and adverse reproductive outcomes in farmer' families-effect of climates condition favouring fungal growth in grains. *Scand Journal of Work Environmental Health*
- [8] Meat inspector manual, (2007) Abattoir Hygiene; Veterinary Public Health National Department of Agriculture Republic of South Africa; edited by Maja M. p 1-101

- [9] Nesbaken T, G Kapperud, DA Caugant (1996). Pathways of Lysteriamonocitogenes contamination in the meat processing industry. *Inter J Food Micro*. 31: 161-171.
- [10] Ryan, K. J. and Ray, C.G. (2004) Sherris Medical Microbiology (4<sup>th</sup>ed.) McGraw Hill. Pp112
- [11] www.wikipedia/slaughterhouse.com
- [12] www. wikipedia.com/Escherchia