PREVALENCE OF *LISTERIA* SPP. IN A CHICKEN CUTTING AND DEBONING ROOM IN A REFRIGERATED SLAUGHTERHOUSE

Tatiana Aparecida Julião¹, Ana Maria Centola Vidal-Martins², Teresa Cristina Lopes Fernandes Garcia³, Adriana Cássia de Oliveira³, Ana Laís Freitas Huet de Oliveira²; Júlio Cesar de Carvalho Baleiro⁴

¹ College of Agriculture Sciences and Veterinary, São Paulo State University, FCAV-UNESP, Jaboticabal, Sao Paulo, Brazil

² College of Animal Science and Food Engineering, University of São Paulo, FZEA-USP, Pirassununga, Sao Paulo, Brazil

³ Veterinary Medica, Ministry of Agriculture Livestock and Supply, Federal Inspection Service, Sao Paulo, Brazil
⁴ College of Veterinary Medicine and Animal Science, University of São Paulo, FMVZ-USP, Pirassununga, Sao Paulo, Brazil

*anamartins@usp.br

Abstract - Chicken meat importing countries have been stimulating Brazil to meet their different microbiological requirements, one of the requirements being the quality of products of animal origin. Listeria monocytogenes has become a major food-transmitted pathogen, causing worldwide concern for the high mortality rate, beyond being easily isolated from chicken meat. The present study's objective was to verify the prevalence of *Listeria* spp. on meat convever belt mats' surfaces in refrigerated poultry slaughterhouses. To this purpose, 160 samples were collected, of which 80 chicken meat cuts and 80 small towels s in an area of 100 cm2 of their respective conveyer belt mats. Prevalence of Listeria spp was found on 88,7% (71/80) of meat cuts and on 49% of conveyer belt mats; highlighting the importance of efficient measures to control such a microorganism in refrigerated slaughterhouses, since current measures are insufficient to insure the absence of the pathogen.

I. INTRODUCTION

Interest in the occurrence of Listeria in food (particularly in Listeria monocytogenes) grew quickly when a series of food-transmitted listeriosis outbreaks occurred in the 1980's [1]. In spite of recent years' technological progress, chicken meat is still contaminated by several microorganisms [2]. Among foods of animal origin that transmit Listeria spp., poultry and its products deserve special attention from researchers due to the association made and between the poultry possible contamination during processing, bringing on the contamination of the final products [3].

Knowing the importance of this microorganism in public health and how it is

considered a sanitary hindrance for meat exportation, the objective of the present study was to evaluate the prevalence of *Listeria* spp. in chicken cutting and deboning rooms in refrigerated poultry slaughterhouses, which are classified as clean areas; from sample meat cuts and the surface of conveyer belt mats.

II. MATERIALS AND METHODS

The samples were collected in cutting and deboning rooms (clean areas air conditioned to 12°C) in chicken freezer-slaughterhouses; submitted to the Service of Federal Inspection (SIF), and to continuous self-control programs (PPHO, HACCP, GMP) with facilities for slaughter, deboning, packing and freezing of the meat cuts. An average160 thousand birds are slaughtered per day, the average speed of slaughtering is 160 birds per minute and products are destined for the domestic market and exportation. The 160 samples were collected during 8 months, with 1 visit a month, totaling 8 visits; 80 of these samples were frozen chicken meat cuts and 80 were from the surfaces of conveyer belt mats.

The cut samples were collected and conditioned in sterile bags and the conveyer belt mat ones were collected through the use of sterile templates (100 cm2) and small towels that were later immersed in tubes containing Listeria Enrichment Broth (LEB), conditioned in isotherm boxes and brought to the LMSA&SA of the University of São Paulo (USP), where they were processed according to a method described by the International Commission on Microbiological Specifications for Foods [4]. The samples of the meat cuts were removed from a 25 gram aliquot and added to 225 mL of LEB; as for surface analysis, the samples were added to 90mL of LEB and incubated at 30°C for 24 hours. Later, 0,1mL aliquots of each sample were transferred to tubes containing 10mL of Fraser broth, incubated at 35°C for 24 to 48 hours. Following the appearance of growth, a sample was transferred to sheets containing oxford agar at 35°C for 24 to 48 hours. The presence of typical Listeria spp. colonies proved positive when the sheets presented pigmented growth with black halos in the agar; their morphology and pigmentation were evaluated using gram coloration and their biochemical features were confirmed.

III. RESULTS AND DISCUSSION

Listeria spp. was isolated in 71 of 80 of the chicken cut samples and in 36 of 80 conveyer belt mat samples, respectively (Figure 1). The technological refrigeration process of the carcasses at an ambient 12°C, manipulation and contact of the carcasses with different sections and equipment of the slaughtering apparatus, collaborate with the evidence that *Listeria* spp. becomes more prevalent in cold products and atmospheres, such as the cutting room, due to *Listeria*'s ability to multiply in cool temperatures, cuts and/or carcasses whose competitive microbiota was reduced during the refrigeration process in chillers [5].



■ Cuts poultry meat ■ Conductive mats of meat

Figure 1. Observed prevalence of *Listeria* spp. in chicken cuts and the surface of conveyer belt mats.

Although L. monocytogenes is a pathogen for humans, all species of Listeria present similar

growth and behaviors. Therefore, the presence of *Listeria* spp. in 88.7% of the cut samples presently observed show the probability of *L. monocytogenes* occurrence, constituting a risk factor in food safety [6].

In a poultry processing plant in the south of Brazil, *L. monocytogenes* has been found in 35.6% of the cut samples, reaching 100% and 93.3% positivity for chests and frozen wings, respectively. The authors proved a larger prevalence of the pathogen than is currently verified in refrigerated meat [7].

The high prevalence of *Listeria* spp. in chicken cuts in the present study was even higher than the prevalence reported by a study in Iran where it was shown that 134 (33.3%) out of 402 poultry product samples were contaminated with *Listeria* spp., with 34.7% of prevalence in raw products and the same was observed in 30.7% in processed products, showing that the processing did not reduce the microorganism's prevalence in the food. [8]

Investigating the presence of *Listeria* spp. in 3685 raw samples (meat, milk and chicken) from northern Spain, and processed products (cured and cooked meat, frozen vegetables and smoked salmon); the largest occurrence of *Listeria* spp. was found in raw chicken meat samples (76.3%), followed by of ground bovine and swine meat samples (62.3%) [9], emphasizing the importance of chicken meat research.

In studies where two chicken processing industries inside São Paulo state were assessed, thev found 23.4% of microorganism contamination on surfaces that touch the food, and 16.8% of surfaces that don't touch the food. This differs from the results of the present study where the prevalence of the microorganism was found to be 49% on surfaces that touch the food [10]. L. monocytogenes presents a high capacity of surface colonization and impermeable biofilm formation, it settles inside plants and, when food is processed, it increases the probability of crossed contaminations and environment contaminations [11].

If pathogen bacteria such as *Listeria monocytogenes* form biofilms, consumers' health may be put at risk, this situation being

particularly preoccupying in industries, which can be in fact justified by the prevalence of *Listeria* spp. in 49% of conveyer belt mat samples of this study [12].

IV. CONCLUSION

Due to high prevalence of *Listeria* spp. mainly found in the cuts and mats in the deboning zone, the importance of a more efficient control of this microorganism in refrigerated slaughterhouses is emphasized; given its dissemination and biofilm formation, and transmission by the food, potentially causing food poisoning.

REFERENCES

- Norton, D. M.; Braden, C. R. *Listeria*, listeriosis, and food safety. New York: Marcel Dekker, 3^a. ed, cap. 10, p. 305-306, 2007.
- Almeida Filho, E. S.; Sigarini, C. O.; Borges, N. F.; Delmontes, E. C.; Ozaki, A. S.; Souza, L. C. Pesquisa de Salmonella ssp. em carcaças de frango (Gallus gallus), comercializadas em feira livre ou em supermercado no município de Cuiabá, MT, Brasil. Higiene Alimentar, v. 17, n. 110, p. 75-76, 2003.
- Dias, M. A. D. Persistência de cepas de Listeria monocytogenes em linha de abate industrial de frango em um matadouro localizado no estado de São Paulo. 2008.
 57 f. Dissertação (Mestrado), Faculdade de Ciências Farmacêuticas, Universidade de São Paulo, São Paulo, 2008.
- International Commission on Microbiological Specifications for Foods (ICMS). Microorganisms in foods: their significance and methods of enumerations.
 ed. University Toronto Press, 1978. 434p.
- Cerutti, M. Programa de qualidade higiênica na produção avícola. In: Berchieri Júnior, A.; Silva, E. N.; Di Fábio, J.; Sesti, L.; Zuanase, M. A. F. Doenças das aves. 2. ed. Campinas: Fundação APINCO de Ciência e Tecnologia Avícolas, 2009. cap. 1.4, p. 53-74.
- 6. Coelho, C. P; Gomide, L. A. de M.; Vanetti, M. C. D.; Passos, F. J. V.; Borges,

M. F.; Siqueira, R. C. S. Efeito, *in vitro*, de pH e nitrito de sódio sobre *Listeria* spp. In: Congresso Brasileiro de Ciência e tecnologia de Alimentos, 16., 1998, Rio de Janeiro, RJ. Anais. Rio de Janeiro: SBCTA, 1998. v.3, p. 889-892.

- Reiter, M. G. R.; Bueno, C. M. M.; Lopez, C.; Jordano, R. Occurrence of *Campylobacter* and *Listeria monocytogenes* in poultry processing plant. Journal of Food Protection, Des Moines, v. 68, n. 9, p. 1903-1906, 2005.
- Fallah, A. A.; Saei-Dehkordi, S. S.; Rahnama, M.; Tahmasby, H.; Mahzounieh, M. Prevalence and antimicrobial resistance patterns of *Listeria* species isolated from poultry products marketed in Iran. Journal of Food Control, v.28, p. 327-332, 2012.
- 9. Vitas, A. I.; Aguado, V.; Garcia-Jalon, I. Occurrence of *Listeria monocytogenes* in fresh and processed foods in Navarra (Spain). International Journal of Food Microbiology, Madison, v. 90, p. 349- 356, 2004.
- Chiarini, E. *Listeria monocytogenes* em matadouros de aves: marcadores sorológicos e genéticos no monitoramento de sua disseminação. São Paulo, 2007. p. 141 Tese (Doutorado em Ciência dos Alimentos) - Universidade de São Paulo -USP.
- 11. Jeong, D.K.; Frank, J.F. Growth of *Listeria monocytogenes* at 100C in biofilms with microorganisms isolated from meat and dairy processing environments. Journal of Food Protection, v.57, p.576-586, 1994.
- 12. Chavant, P.; Martinie, B., Meylheuc, T.; Bellon-Fontaine, M. N.; Hebraud, M. *Listeria monocytogenes* LO28: Surface Physicochemical Properties and Ability To Form Biofilms at Different Temperatures and Growth Phases. Applied and Environmental Microbiology, Washington, D.C., v.68, n. 2, p. 728-737, fevereiro 2002.