

Caridad Valladares

Manuel Roca

Food Industry Research Institute, Havana, Cuba

Siomara Jares

## INTRODUCTION

The microbial contamination of meat during primary processing is of paramount importance to its overall quality. Contamination levels have both a quantitative importance- counts ranging normally between  $10^4$  cells  $\text{cm}^{-2}$  and  $10^6$  cell  $\text{cm}^{-2}$  for normally good and deficient hygienic practices- and a qualitative one, as several species of potentially dangerous pathogenic microorganisms may be present.

Microbial growth, usually hindered by refrigeration, must be carefully considered in the hot boning situation, where cutting and handling smear bacteria all over the several-fold extended meat surface.

This paper deals with microbiological results obtained in hot deboning trials on a pilot plant scale, in order to assess the feasibility of this technology in Cuban sub-tropical conditions.

## MATERIALS AND METHODS

Slaughtering was carried out according to industrial standards. Primary operations included captive-bolt stunning and mechanical dehiding. One side of each of 4

cattle slaughtered was immediately debone and the hot meat packed into plastic trays and corrugated cardboard cartons covered in either case with polyethylene film and chilled in air at  $4^\circ\text{C}$  and  $1 \text{ ms}^{-2}$  air velocity.

The four remaining sides were conventionally chilled, deboned after 24 hours refrigeration, the meat hung in stainless steel trees and cloth covered, as usual under commercial conditions in Cuba.

A transport stage was also simulated, letting the refrigerated meat stand in air at  $28^\circ - 30^\circ$  for 5 hours, which would be equivalent to transport in a closed non-refrigerated truck or van, not uncommon in our conditions.

Samples for microbiological analysis were taken using the swabbing technique of Kitchell *et al* (1). Total counts for mesophiles and psicrotrophs were obtained on PCA, incubating at  $30^\circ\text{C}$ , 48 hr. and  $4^\circ\text{C}$  for 7 days, respectively. Coliform counts were obtained on Red violet bile agar, at  $37^\circ\text{C}$ , 20-24 hr. Standard dilution plate counts were made in all cases.

## RESULTS AND DISCUSSION

Table 1 shows typical microbial counts in normal industrial operation in Cuban slaughterhouses. These results indicate quite acceptable hygienic conditions even by literature standards (2,3,4), and so hot deboning was considered potentially feasible, at least from the point of view of primary operations. Similar conclusions could be derived from data for refrigerated carcasses ( Table 2).

Sampling point	Total viable mesophile count	Total psicrotroph count	Coliform count
Collar	4,08	2,42	2,62
Diaphragm	4,02	2,33	2,44
Leg	3,92	2,72	2,10
Side average	4,01	2,49	2,39

Table 1- Microbial counts on recently slaughtered beef sides. Average  $\log_{10}$  values

Sampling point	Total viable mesophile count	Total psicrotroph count	Coliform count
Collar	4,33	2,66	2,69
Diaphragm	3,69	2,96	2,17
Leg	4,21	2,58	2,52
Side average	4,08	2,57	2,43

Table 2- Microbial counts of refrigerated beef sides, 24 hours post mortem. Average log<sub>10</sub> values.

In preliminary hot deboning trials ( Table 3 ) unacceptably high counts were obtained for hot deboned meat in packaging options. Cardboards cartons were eliminated, since core temperature could not be brought down as rapidly as required, which is an essential (5).

Sampling point	Total viable mesophile count	Total psicrotroph count	Coliform count	Deboning System
Collar	4,04	2,00	2,17	Conventional
Diaphragm	3,69	1,00	2,07	
Leg	3,77	2,37	1,00	
Side average	3,83	1,78	1,74	
Plastic tray	6,72	5,96	4,47	Hot
Cardboard carton	8,72	7,77	5,47	

Table 3- Hot vs conventional deboning. Preliminary results. Average log<sub>10</sub> values of 4 replicates.

The plastic tray option was given a second trial, taking special care regards hand-washing and utensil disinfection. Results are present in Table 4, showing the effectiveness of stringent requirements for hygienic operation. Results are good, even

after the abuse of simulated transport.

Sampling point	Total viable mesophile count	Total psicrotroph count	Coliform count	Deboning System
After deboning	3,16	2,83	1,61	Hot
After refrigeration	4,33	4,52	2,82	
After "transport"	4,98	4,81	3,37	
After deboning	3,83	3,69	1,39	Conventional
After "transport"	4,83	5,44	2,85	

Table 4- Hot vs conventional deboning. Average log<sub>10</sub> values of 4 replicates.

#### ACKNOWLEDGEMENT

Thanks are due to Dr Charles L. Cutting for helpful advise.

## REFERENCES

- 1- Kitchell, A. G; Ingram, M; Hudson, W. R. (1973). Sampling microbiological monitoring of environments. Academic Press, London. New York.
- 2- Ingram, M; Roberts, T. A (1976). The microbiology of the red meat carcass and the slaughterhouse. Roy. Soc. Hlth. J. 96 (6) 270.
- 3- Roberts, T. A (1980). Contamination of meat. The effects of slaughter practices on the bacteriology of the red meat carcass. Roy. Soc. Hlth. J(2).
- 4- Taylor, A.A; Show, B. C; (1980). Meat deboning beef with and without electrical stimulation. Meat Science (5) 109-123
- 5- Herbert, L.S; Smith, M.G (1980). Hot boning of meat. Refrigeration requirements to meat microbiological demands. CSIRO. Food Press. Vol 40 3/4