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## Microbiological quality of mechanically recovered meat made from pork and poultry

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## Background/ Objectives

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The quality of mechanically recovered meat, its good technological characteristics and comparatively low cost make the product a profitable and useful raw material.

Meat removed from bones by machines is generally termed mechanically deboned meat (MDM), mechanically deboned tissue (MDT), mechanically separated meat/ tissue (MSM/ MST) or mechanically recovered meat (MRM).

Both red meat and poultry meat can be a source of pathogens such as Salmonella spp., Campylobacter spp., enterohaemorrhagic E. coli O157:H7, Listeria monocytogenes, Yersinia enterocolitica, Staphylococcus aureus etc., as well as spoilage bacteria, such as Pseudomonas, responsible for the development of rancidity. Therefore it is essential that all sanitary and hygienic measures concerning the raw material te be used for the production of MRM are absolutely restricted and controlled. Meat quality can be judged very well by the quality of microbes (NURMI, 1997). In the present investigation, the microbial quality of MRM was investigated.

### Material and methods

#### MRM meat production

During the period February 1 to July 1, 1997, 36 samples of two types of MRM produced with the "Lima" and "Protecon" separator machines, respectively, were investigated for pathogenic bacteria.

For the detection of Salmonella, Campylobacter, Yersinia, enterobacteriaceae and total cell counts, methods were used as set down in paragraph 35 of the LMBG and in ISO 6579 : 1993 (E), ISO 10272 : 1995 and ISO 10273. Listeria were cultivated in a Listeria enrichment broth (Oxoid SR 149) at 30° C for 24 h. For the detection of Listeria, two selective agars (Palkam and Oxford agar [Oxoid]) were used, with cultivation tor 24 h, 48 h, and 7 d, at 37° C. All detected bacteria were further differentiated serologically or biochemically.

#### Results

The results of bacteriological investigations of "Protecon" type MRM are presented in Table 1.

Further differentiation of the salmonella provided confirmation of the presence of S. typhimurium and S. dublin, while the Yersinia were proved to be Y. enterocolitica Serovar 0:5, Biovar 1a (non-pathogenic microbe). Biochemical differentiation of Listeria demonstrated L. grayi, L. welshimeri, and L. innocua. When host animal species are compared, C. coli is found most frequently in pigs and pork. The following Campylobacter were detected: C. coli, C. sputorum faecalis and C. jejuni II.

#### Discussion

MRM - obtained from pork and in recent years also from poultry meat - is still used for the production of boiled sausages and meatballs. The hygienic quality of these products in our investigations, a large part of these products proved to be of poor hygienic quality.

MRM can show a bacterial content as high as  $1.5 \times 10^3$  to  $3.4 \times 10^6$  tvc/g for total plate count, and  $1.8 \times 10^4$  to  $6.5 \times 10^6$  tvc/g for enterobacteriaceae. The rates of detection of Salmonella (16.6%), Campylobacter (50.0%) and Listeria (22.2%) are also on an elevated level.

The results of investigation in other countries also show a high rate of contamination of MRM with salmonella and Staphylococcus aureus, as well as with other coliform bacteria (BANKS and BOARD, 1993; ADAMS and MEAD, 1983). The importance of the degree of freshness of the original material cannot be overestimated.

An expert group of the Scientific Veterinary Committee, which advises the EU Commission in matters concerning public health, has recommended the maximum bacterial contamination of separated meat shown in Table 2.

The samples we investigated were obtained immediately after production and transported chilled to the laboratory within three hours. If MRM is not cooled immediately, the total plate count can soon exceed the barrier to spoilage levels. When raw materials with high contamination levels - resulting in MRM with high contamination levels - are used, it is not possible to safely guarantee the criteria of quality products even in heated products such as boiled sausages, due e.g to heatstable toxins of some bacteria



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<sup>Code</sup> of Practice for MRM and Poultry Meat (1994): <sup>Recommended</sup> International Code of Practice for the Production, Storage and Composition of MRM... (CAC/RCP 32-1983) <sup>Codex</sup> Alimentarius, Volume 10 - 1994

<sup>Council</sup> Directive 6414331E~C, last amended and updated by Directive 95123 of June 1995 Article 2c

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# Tables

Table 1: Presence of pathogenic bacteria in "Protecon" type MRM

No. samples <sup>investigated</sup>	total plate count	Enterobacteriaceae	Salmonella (%)	Campylobacter (%)	Yersinia (%)	Listeria (%)
		$\frac{1.8 \times 10^4}{\text{to } 6.5 \times 10^6}$	16.6	50.0	5.5	22.2
<sup>36</sup> from poultry	$4.2 \ge 10^5$	$2.7 \times 10^7$	32.3	38.5	-	11.6

Table 2: Recommended bacteriological guidelines for MRM (NURMI et al. 1997)

	total plate count 25° C/ 3 d (ISO)		Enterococci (Slanetz and Bartley Medium) 44° C/ 2 d		Enterobacteriaceae	
	m	М	m	М	m	М
MRM red meat	$5 \ge 10^5$	5 x 10 <sup>6</sup>	$5 \times 10^3$	5 x 10 <sup>4</sup>	$5 \times 10^3$	5 x 10 <sup>4</sup>
MRM poultry meat	$5 \ge 10^5$	5 x 10 <sup>6</sup>	5 x 10 <sup>3</sup>	5 x 10 <sup>4</sup>	5 x 10 <sup>4</sup>	5 x 10 <sup>5</sup>