6:30

Effect of sulphamethazine on the ripening of Italian "salame casareccio"

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<u>SUMMARY</u>: Few differences in the chemical composition and in the sensorial evaluation have been detected among five groups of country-style salamis (salame casareccio) whether with or with sulphamethazine added at different concentrations. These might be due to different diameters of the casings and consequently to the slightly different rate of dehydration and within the groups and within the groups.

Only slight differences have been recorded in the total aerobic microbial count, Enterior bacteriaceae and Lactobacillaceae among the groups during the ripening period. On the other hand a significative difference in the other significative difference in the significative differe hand a significative difference in the growth of Micrococcaceae has been detected between the group of salamis with the higher concentration of sulphamethazine and the other groups the This difference was even more evident in the Micrococcaceae/TMC ratio which was less the 1°/.. until the end of the 8-week ripening period in the group with the higher sulphamethor h^{0} concentration (0.5 ppm). The two controls without sulphamethazine and the group with of this compound added showed a maximum Micrococcaceae/TMC ratio at the fourth week. Although Micrococcaceae is regarded an important family of bacteria in ripened meat products, no marked differences were found in these country-style salamis, probably because the short ripening period required made the metabolic satisfies period required made the metabolic activities of Micrococcaceae less relevant.

INTRODUCTION: Meat inspection surveys frequently detect sulphonamide residues in surveys frequently detect sulphonamide residues in such as the subtissue due to their wide use in pig husbandry, both for therapeutic purposes and for prevention of common diseases. Subbangride tion of common diseases. Sulphonamides must be withdrawn 21 days before slaughtering will administered at 3750-5000 mg/Kg in fact for administered at 3750-5000 mg/Kg in feed for 3-5 days because they are slowly metabolized Contamination of non medicated feed occurs when medicated feed is prepared at the mill, recycling of sulphonomides may occur then it recycling of sulphonamides may occur through residues in faeces and urine (Epstein et 1988). 1988).

Italian researchers have recently reported that 10.5% of 123 aged cured Italian hamble bought in retail stores in several regions of our country, contained over 0.1 ppm suppr methazine residue (Cortesi et al 1000) of our country, contained over 0.1 ppm suppr methazine residue (Cortesi et al.,1990). Moreover, 14.6% of 48 aged cured hams processed abroad were positive for sulphamethazine and the sulphamethaz abroad were positive for sulphamethazine residue at 0.1 ppm (Cattaneo et al., 1990). A monthage toring programme set up in central train (Mither at 0.1 ppm (Cattaneo et al., 1990). A monthage (Mither at 0.1 ppm (Cattaneo et al., 1990). toring programme set up in central Italy (Umbria region) led to the conclusion that 18 4.11% of regularly slaughtered pigs in 1989 had residues of antimicrobial compounds, inhibition effect in the microbial test used to detect residues was positive in 30% of covering only when trimethoprim was added. Suggesting the only when trimethoprim was added, suggesting the presence of sulphonamide residues (Severiment et al., 1990). Since the sensitivity of the test et al.,1990). Since the sensitivity of the test used to sulphonomides is about 0.5 ppm, percentage of pigs with a lower amount of these residues might even be higher.

This experiment was designed to evaluate whether and how sulphamethazine (SMT), is one of the most widely used sulphonamides in pig husbandry, affects the quality of locity ripened salami. In the microbiological evaluation particular emphasis was given to Latter bacillaceae, Micrococcaceae and Enterobactorizecoust bacillaceae, Micrococcaceae and Enterobacteriaceae since their growth is of utmost important to the ripening of Italian salami (Cantoni et al., 1989; Censi et al., 1989).

MATERIALS and METHODS: Five groups of Italian "salame casareccio" were prepared by mixing - the first group was added with 100 ml of sterile distilled water and was used as a control

^{In diameter.} All salamis at the end of the ripening were classified as good quality products by Means of a sensorial evaluation. Colour, smell and flavour were typical of this kind of or^{ja} ^{pr}oduct, likewise its aspect and consistency.

Data on the total mesophylic aerobic count (TMC) are reported in Figure 1. The values r_{unged} from 695*10³ to 3.5*10⁹ ufc/g and the maximum values were reached during the last Weeks of ripening. The growth of mesophylic aerobic bacteria was similar for all groups, except for the first which had three times the amount of the other groups during the first

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^{Data} on Lactobacillaceae count are given in Figure 2. The proliferation of this bacteria ^{Nos} so fast that near maximum values were reached in all groups in the first week of ripening. Values of Micrococcaceae at given times are reported in Figure 3. Until the fifth week the amount of this bacteria in the group with the highest concentration of sulphamethazine (0.5 ppm) was less than 4*10⁴ ufc/g, reaching a value of half a million ufc/g only at the ^{eighth} week. The two controls had a different increase in the number of Micrococcaceae, with ^{Noximum} Values (4.75*10⁵; 4.3*10⁵) at the fourth week. The proliferation in the groups with 0.05 and 0.1 ppm sulphamethazine was higher than in the group with 0.5 ppm and lower than the the ^{control} groups. The difference in ratio between Micrococcaceae and total mesophylic aeroblc bacteria is even more explicative (Fig. 4). In the group with the higher SMT concentration (0.5 PDm) the ratio was less than 1°/... until the end of the 8-week ripening period, where 10^{10} the the two controls without SMT and the group with 0.05 ppm SMT the maximum ratio (>2°/...)was thoserved as early as the fourth week. The group with 0.1 ppm SMT had a lower proliferation than the controls and the group with 0.05 ppm during the first five weeks, reaching a maximum ^{ratio} at the sixth week.

the second group was added with 100ml of sterile NaOH 0.001N and was used as a control for t for the SMT solvent; the third group was added with 100ml of sterile sulphamethazine/NaOH solution (0.5%p/v);

the fourth group was added with 100ml of sterile sulphamethazine/NaOH solution (1%p/v); the sulphamethazine/NaOH solution (5%p/v).

the fifth group was added with 100ml of sterile sulphamethazine/NaOH solution (5%p/v). The final concentrations of sulphamethazine in the products were therefore 0.05 ppm, 0.1 ppm Ond D r ^{and} 0,5 ppm, respectively. An even spread of cure ingredients and sulphamethazine was achieved by Mixing Mixing the put into natural casings and ripened by mixing accurately. Then the five types of mixture were put into natural casings and ripened accurately. Inc.

Protein content, moisture, pH and Aw were evaluated in cured ground meat before processing in and in salamis taken from each group. The sensorial quality of the salamis was evaluated by a Doperations were performed according to current by a ^{Danel} test. The following microbiological evaluations were performed according to current ^{Methode} Methods: total aerobic microbial count (Tryptone agar media-BBL); Lactobacillaceae (LBS agar-BBL); Mi BBL); Micrococcaceae (MSA agar-BBL); Enterobacteriaceae (Violet Red Bile Dextrose agar-Oxoid); Staphylo Staphylococcus aureus (Baird-Parker medium-Oxoid); Enterococci (Barnes medium); Coliformis bacterio bacteria (Brilliant green bile broth 2%-BBL); sulphite-reducer Clostridia (Brain heart infu-^{\$10}h 0000 ^{\$1}on agar-BBL).

RESULTS and DISCUSSION: Results of the chemical analysis are reported in Table 1. The values of the first two weeks of ripening, ^{then} rose in 5.43 to 5.72. No relevant differences then rose to slightly higher final values ranging from 5.43 to 5.72. No relevant differences ^{mong} to slightly higher final values ranging from 5.45 to 5.72, no rote kind the groups were observed. Moisture and protein content varied, as is usual in this of kind of Country-style product, according to the dehydration rate and thus the differences

6:30

Enterobacteriaceae showed very little proliferation in all the groups except the first off which reached a value of 3200 ufc/g in the third week. Any effect on the growth of $Enter^{\ell}$ bacteriaceae could be attributed to the addition of sulphamethazine.

The hygienic quality of the products was good: coagulase-positive Staphylococcus aureus was never detected; Enterococci were constantly at low levels (500-260000 ufc/g); Colifornis bacteria were less than 0.3 ufc/g; sulphite-reducing Clostridia were absent.

<u>CONCLUSIONS</u>: The addition of 0.5 ppm SMT to pigmeat used in manufacturing "salame casare" cio" caused a significant difference only in the growth of Micrococcaceae and when the con centration of SMT was 0.5 ppm. The quality of the ripened salamis was not affected, perhod because the metabolic activities of this group of bacteria are not of such importance these salamis with a short ripening period.

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PHYSI()-(C)	HEM1	CAL	EV	ALUAT	ION	OF	FIVE	GROUPS	OF	"SALAME	CASABECCIO	p
ADDED	OR	NOT	WIT	Н	SULPH	AMET	HA.	ZINE	(SMT).			Sar 7 7 mer 7 77 V han hear too? als too	

		time (weeks)									
		0	1	2	3	4	5	6	8		
	groups*										
	1	5,91	5,39	4,97	5,25	5,23	5,38	5.32	5.50		
	2	5,91	5,40	5,48	5,63	5,34	5.54	5.56	5.48		
На	3	5,91	5,33	5,22	5,58	5,48	5.62	5.54	5,50		
	4	5,91	5,36	5,22	5,48	5,34	5.36	5.56	5.72		
	5	5,91	5,32	5,29	5,50	5,47	5,38	5,54	5,43		
MOISTURE%	1	61,90	49,49	47.55	42.60	40.10	36.56	38.48	74 D7		
	2	61,90	48,10	47,37	40.51	40.04	39.40	34.67	30.73		
	3	61,90	49,39	45.64	44.97	39.02	40.30	37.23	37 10		
	4	61,90	49,10	46,36	41,37	37.84	41.70	33.41	33.83		
	5	61,90	48,52	48,18	40,21	37,84	41,26	32,10	36,99		
	1	n.d.	n.d.	0,94	0,93	0,91	0.89	0.87	0.87		
	2	n.d.	n.d.	0,93	0,91	0.90	0.88	0.87	0.88		
AW	3	n.d.	n.d.	0,93	0,92	0,91	0.89	0.88	0.86		
	4	n.d.	n.d.	0,93	0,92	0.90	0.87	0.86	0.84		
	E.J	n.d.	n.d.	0,93	0,92	0,90	0,88	0,89	0,86		
	1	n.d.	19,30	17,79	19.27	n.d.	n.d.	19.51	21.15		
	2	n.d.	18,45	18,87	22,44	n.d.	n.d.	22.16	24.17		
PROTEIN%	3	n.d.	19,65	18,39	20,23	n.d.	n.d.	22.90	23.08		
	4	n.d.	17,85	18,46	20,11	n.d.	n.d.	22.31	21.23		
	5	n.d.	17,94	17,96	19,85	n.d.	n.d.	19.97	19.17		
									an e g de s		

*1=control; 2=NaOH control; 3=SMT 0.05ppm; 4=SMT 0.1ppm; 5=SMT 0.5pp





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