g of Auctural and Biochemical Changes During Aging of Hot Deboned and Electrically Stimulated Bovine Muscles

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#### SUMMARY tori

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thect of electrical stimulation on the structural changes and condition of myofibrillar proteins from hot deboned bovine Wecles was investigated during postmortal aging.

The muscles were excised from both halves of warm carcasses, 1 hour p.m.: M. longissimus dorsi (LD), M. biceps femories <sup>(Huscles</sup> were excised from both halves of warm carcasses, in the providence of the Wrapped in PVC foil and aged at 0°C. Tenderness - sensory and instrumentally and solubility of total proteins were Refermined in all muscles. The content of free amino-acids and the ultrastructural changes were determined only in SM: 1, 8, <sup>{|</sup> and 42 days after aging.

The obtained results show that ES prevents cold shortening in LD and SM, while in BF it was not estimated neither in ES nor <sup>th</sup> <sup>muscles</sup>. The tenderness of NS muscles reached between 21<sup>st</sup> and 42<sup>nd</sup> day, was achieved in ES muscles between 8<sup>th</sup> and 21st day.

The increase of tenderness during the whole aging period was accompanied by protein solubility increase til the 8<sup>th</sup> e.g. 21<sup>st</sup> <sup>Ay, and</sup> after that by decrease in both groups. In the same time, the cross-striation in ES muscles disappeared, and in NS uscles the zig-zag form of the Z-membrane was observed.

## MIRODUCTION

<sup>Ne</sup> it is well known, tenderness of bovine muscles increases during aging. However, aging is a long process, so the way how <sup>b</sup> shorten it is permanently investigated.

<sup>electric</sup>ally stimulated muscles pH droppes in a short time to the value optimal for the activity of proteolytic enzymes. So tivity of these enzymes is expected to start earlier, and in that way, the aging e.g. tenderization of the meat can be <sup>tor these</sup> enzymes is expected to start earlier, and in that thay, the second start gravely and the second start earlier, and in that thay, the second start earlier and the second start earlier and in that thay, the second start earlier earlier

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here are very few data available on the influence of electrical stimulation on the characteristics of hot-deboned muscles during aging. So the aim of this work was to investigate this influence.

# MATERIALS AND METHODS

Were been seed for the investigation. Stunning and bleeding <sup>vere</sup> <sup>Performed</sup> in the usual way. After the treatment on the line, before weighing and final washing, approximately 35-40 min. M. Semi-M<sub>semimembranosus</sub> (SM) and M. biceps femoris (BF).

Muscles from the right halves were electrically stimulated (ES) (after excising) for 120 sec. with 14 Hz pulses, with constant Muscles from the right halves were electrically stimulated (ES) (after excising) for 120 sec. with 14 Hz pulses, with constant of the right halves were electrically stimulated (ES) (after excising) for 120 sec. with 14 Hz pulses, with constant the right halves were electrically stimulated (ES) (after excising) for 120 sec. with 14 Hz pulses, with constant the right halves were electrically stimulated (ES) (after excising) for 120 sec. with 14 Hz pulses, with constant the right halves were electrically stimulated (ES) (after excising) for 120 sec. With 14 Hz pulses, with constant the right halves were electrically stimulated (ES) (after excising) for 120 sec. With 14 Hz pulses, with constant the right halves were electrically stimulated (ES) (after excising) for 120 sec. With 14 Hz pulses, with constant the right halves were electrically stimulated (ES) (after excising) for 120 sec. With 14 Hz pulses, with constant the right halves were electrically stimulated (ES) (after excising) for 120 sec. With 14 Hz pulses, with constant the right halves were electrically stimulated (ES) (after excising) for 120 sec. With 14 Hz pulses, with constant the right halves were electrically stimulated (ES) (after excising) for 120 sec. With 14 Hz pulses, with constant the right halves were electrically stimulated (ES) (after excising) for 120 sec. With 14 Hz pulses, with constant the right halves were electrically stimulated (ES) (after excising) for 120 sec. With 14 Hz pulses, with constant the right halves were electrically stimulated (ES) (after excising) for 120 sec. With 14 Hz pulses, with constant the right halves were electrically stimulated (ES) (after excising) for 120 sec. With 14 Hz pulses, with constant the right halves were electrically stimulated (ES) (after excising) for 120 sec. With 14 Hz pulses, with constant the right halves were electrically stimulated (ES) (after excising) for 120 s Meaks from the right halves were electrically stimulated (ES) (after excising) for 120 sec. with the right halves were electrically stimulated (ES) (after excising) for 120 sec. with the right halves of 1, 1987). The Muscles of 32 for 5 msec and pauses of 70 msec - using a device of our construction (Petrović Ljiljana et al, 1987). The  $\eta_{\text{USCles from the left halves were not stimulated (NS).}$ 

<sup>0</sup><sup>heg</sup> from the left halves were not stimulated (NS). <sup>0</sup> to p.m. all excised muscles were cut into 4 pieces, packed in PVC foil under vacuum and kept in the refrigerator at the steaks of the ste <sup>110</sup> to 5.°C till the moment of investigation. The samples were investigated after 1, 8, 21 and 42 days p.m. Two 3 cm thick steaks Were cut from every sample. One steak was used for instrumental investigation of tenderness was determined sensorily in still <sup>NB</sup>. The other cut was treated by grilling to an internal temperature of 55°C. The tenderness was determined sensorily in still hot samples.

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From the remaining piece, a sample was taken (always from the same place) for the investigation of ultrastructure with electron microscop, and after the removing of outside fatty and connective tissue it was ground, homogenized and kept in hermetically closed containers. This sample was used for the determination solubility of total proteins in 1 M KCl pHe<sup>7/2</sup> (Awad et al, 1968) and content of free amino-acids (only SM was used).

Six samples from both groups (NS and ES) were investigated, and the results were statistically analyzed (Hadživuković, <sup>1984</sup> using analysis of variance.

#### **RESULTS AND DISCUSSION**

The results obtained during our investigations are presented in 4 graphs, 1 table and 2 pictures.





Graph. 1. Tenderness (WB) in three muscles excised early p.m. (NS and ES), during aging (n = 6)

Graph 2. Sensorily evaluated tenderness in three  $m^{USCRS}$  excised early p.m. (NS and ES), during aging (n = 6)

The first day p.m. the lowest tenderness was determined in NS muscles: SM (8.65 kg), followed by LD (8.07 kg) and BF (6.93 kg). In the same period, the ES muscles were more tender: LD and SM more (6.53 e.g. 6.29 kg), and BF somewhat less (6.08 kg).Both NS and ES muscles became more tender during aging, and the difference between the tenderness of ES and NS muscles remained till the end of aging. It is important to underline that the tenderness estimated in ES muscles after 21 day of aging was achieved in NS muscles at the end of the aging period e.g. after 42 days.

The influence of muscle kind on the tenderness (WB) is not significant, neither of tested treatments (Table 1). However, the influence of applied process (ES) and aging time is highly significant (P<0.001).

Table 1. Analysis of variance of the influence of muscle kind, treatment and aging time on the investigated characteristics

Source of	Degree of	F-value		
variation	freedom	Tenderness		Solubility of
		WB	sensory	total proteins
Muscle (M)	2	2.25	3.33*	0.28
Treatment (Tr)	1	37.29***	39.48***	1.10
Time (T)	3	56.11***	98.03***	19.08***
Interaction M-Tr	2	1.82	0.62	0.02
Interaction M-T	6	1.37	1.36	1.66
Interaction Tr-T	3	0.89	2.07	0.33
Remainder	126	-	-	-
Total	144	-	2	-

significantly different: \* (P<0.05); \*\* (P<0.01); \*\*\* (P<0.001)

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with the results presented in graph 2 show that the sensorily evaluated tenderness of thermically treated ES muscles is higher one plin by p.m. than the one of NS muscles. It is especially the case with SM., where the tenderness of NS muscle was graded 2.99 <sup>1/2</sup> <sup>Ind</sup> <sup>of</sup> ES muscle 4.22. During aging, all NS and ES muscles became more tender. Though at the end of the aging the ES Were evaluated as more tender than the NS ones, the difference determined the first day p.m. gradually decreased til he 42nd day, especially in the case of LD.

<sup>he</sup> analysis of variance showed (Table 1) that the influence of kind of muscle (P<0.05), and the applied treatment and time of <sup>ang are significant</sup> (P<0.001) for the tenderness evaluated by sensory method, what is not the case with their interaction. Was found that the free amino-acids (graph. 3) was increasing til the end of aging.



<sup>Qraph, 3.</sup> Content of total free amino acids in SM excised Graph 4. The solubility of total proteins of three muscles <sup>om carcasses</sup> early p.m. (NS and ES), during aging (n=6) excissed early p.m. (NS and ES), during aging (n=6)

Between the 21st and 42<sup>nd</sup> day of aging, a more significant increase of free amino-acids content was estimated and it was Igher in ES than in NS muscles

the determination of total proteins solubility resulted in oposite findings (graph. 4). Namely, the solubility increases til the 8th the solubility of total proteins solubility resulted in oposite findings (graph. 4). Namely, the solubility increases til the 8th the solubility of total proteins solubility resulted in oposite findings (graph. 4). Namely, the solubility increases til the 8th the solubility of total proteins solubility resulted in oposite findings (graph. 4). Namely, the solubility increases til the 8th the solubility of total proteins solubility increases til the solubility of total proteins solubility resulted in oposite findings (graph. 4). Namely, the solubility increases til the solubility increases til the solubility increases til the solubility of total proteins solubility increases til the solubility increases till the both in NS and ES muscles: LD and BF, and in SM til the 21<sup>st</sup> day. In all groups of muscles (NS and ES) solubility <sup>tecreased</sup> further on. However, the solubility was somewhat higher in ES muscles.



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<sup>Figure</sup> 1. Electron micrographs of SM muscle after 42 days of aging (x 30.000) a) NS muscle, b) ES muscle

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#### CONCLUSIONS

The results of tenderness determination indicate that eletrical stimulation prevents cold shortening in hot-boned LD and SM.<sup>III</sup> BF this phenomenon was not observed neither in ES nor in NS muscles.

The tenderness estimated in NS muscles between 21st and 42nd day of aging was achieved in ES muscles between 8th and 21st day.

During the whole aging period the content of free amino acids was increasing. The solubility of total proteins increased till the 8th (LD and BF) e.g. 21st day (SM), and after that decreased in both investigated groups.

In the same time, the tenderness increase in ES muscle was accompanied with the loss of cross-striation, and in NS muscle with the zig-zag form of Z-membrane.

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