INTER AND INTRAMUSCULAR VARIATION ON SENSORY CHARACTERISTICS OF BEEF STEAKS

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tenderiess is the most important quality trait for consumer acceptance and the variability in this concept has increasing problem for the retailers. Tenderness is extremely variable between at tenderness is the content of the retailers. Tenderness is extremely variable between carcasses, between muscles are the come muscle, and in part can be due to total colleges content of the content o an increasing production of the same muscle, and in part can be due to total collagen content and its solubility, influencing the soover within the same the consumers of meat (Denoyelle and Lebihan, 2003, Torrescano et al., 2003). Sensory analysis in determines the quality attributes that consumers are going to value. In this sense, Shackelford et al., (1995) determines the distribution of hardness in ten different bovine muscles and found that sensory values in were very variable. The objective of this study was to observe inter and intramuscular variability on sensory were very value of the entire muscles of different regions of the beef carcass included in the label from PGI "Carne de la Sierra de Guadarrama".

the male yearlings from Limousin, Charollais or Brown Swiss bulls, and Avileña-Negra Ibérica dams (a bovine god from the Central area of Spain) were studied. All animals were the quality label from the PGI "Carne de la Sierra b Guadarrama" (a Protected Geographical Indication of Madrid). Animals employed in this study were entire males 13-15 months old, 391 kg HCW and a carcass classification value of R3 (European Union, 1991). They were adlitered in a commercial abattoir and processed according to the rules. M. longissimus dorsi (thoracic portion) (LD), t smitendinosus (ST) and M. supraspinatus (SS) were removed at 7 days of ageing postmortem and cut in steaks (2 thick) of three homogeneous regions (1, 2 and 3). The regions in LD were cranial, middle and caudal, while that in 57 and SS muscles were dorsal, middle and ventral regions, respectively. Attributes measured included pH, water ding capacity, collagen composition, and intramuscular fat content. Also, a compression textural measurement by IPA (texture profile analysis) method was studied (Velasco et al., 2005). One cm x 1 cm-strips were made from each to for texture assessment on raw meat. The texture analysis was made with a cylindrical 10 mm-diameter probe of etonite and with a deformation force of 75%. Sensory analysis was performed on samples from three muscles from each of fifteen animals, grill-cooked till 80°C (internal temperature). Samples were randomly served to a trained twelve ber sensory panel. Sensory analysis evaluated hardness, springiness and juiciness, completed with the parameters of fat sensation (in mouth), flavour intensity, number of chewings until swallowing and overall analysis. Scales used to usess intensity of every parameter were interval scales of 10 cm long. Statistical analyses were carried out using the malysis of variance of the GLM procedure of the Statgraphics Plus (1994). Differences between means were compared

Results and Discussion

aing Duncan test

Florts of different muscles and regions inside these muscles, on pH, water-holding capacity, and intramuscular fat and collagen contents are shown in Table 1.

Table 1: Arithmetic means and mean squares of the error of pH, WHC, intramuscular fat and collagen parameters of seles and regions studied

This cours study	icu.									
	Muscles (M)				Regions (R)					
	LD	ST	SS	Sig	1	2	3	Sig	MxR	MSE
PH	5.53ª	5.54ª	5.65 ^b	*	5.59	5.54	5.59	NS	NS	0.015
WHC (%)	15.59a	18.21 ^b	18.90^{b}	*	18.51	17.24	16.95	NS	NS	9.16
Intramuscular fat (%)	2.04^{ab}	1.42a	2.50^{b}	**	2.07	2.00	1.90	NS	NS	0.82
(mg/g muscle)	6.04 ^a	11.54 ^b	10.39^{b}	***	9.19	9.49	9.28	NS	NS	5.48
(mg/g muscle)	4.77 ^a	9.11 ^b	7.53^{b}	***	7.17	7.03	7.21	NS	NS	3.60
Solubility (% total)	13.07	14.84	12.90	NS	13.67	12.79	14.35	NS	NS	23.90

in the same row not followed by a common letter differ significantly. Sig: Significance; NS (non significance), * (p<0.05), ** (p<0.001), MSE: mean square of the error. LD= M. longissimus dorsi, ST= M. semitendinosus, SS= M. supraspinatus; 1 (cranial in LD, and dorsal in ST and SS), 2 (middle), 3 (caudal in LD, and ventral in ST and SS). WHC (water-holding ent, expressed as percentage of expelled liquid), TC (total collagen), IC (insoluble collagen).

Significant intermuscular variation was detected at 7 days postmortem, with higher pH value in SS than in LD and SS than in ST, while a higher quantity of exudate was seen in ST SS than in LD muscle. Thus it could be due to the fact that the greater the pH the greater the WHC (Purchas, 1990) Total and insoluble collagen contents were higher in ST and SS than in LD as Torrescano et al., (2003) noted No

Table 2: Arithmetic means and mean squares of the error of textural and sensorial parameters of muscles and studied.

	Muscles (M)				Regions (R)				- 2	and regio	
	LD	ST	SS	Sig	1	2	3	Taken 1		_	
TPA Hardness (N)	206.33 ^a	369.96 ^b	302.72 ^b	***	262.19 ^a	251.61	365.20 ^b	Sig	MxR	Ne	
TPA Springiness (g)	74.60°	78.38 ^b	76.80^{ab}	*	74.63	77.14	78.01	**	NS	MS	
TPA Chewiness	4211.9 ^a	11966.6 ^b	8527.5°	***	6879.2ª	7044.2ª	10782.7 ^b	NS	NS	847	
Hardness	4.47 ^a	5.08 ^b	5.06^{b}	*	4.92	4.66	5.03	*	NS	20.	
Springiness	4.42	4.50	4.88	NS	4.80	4.42	4.59	NS	NS	0.2	
Juiciness	2.90^{a}	1.97 ^b	2.42^{ab}	*	2.40	2.37	2.53	NS	NS	0.2	
at sensation	2.56	2.06	2.35	NS	2,23	2.34	2.41	NS	NS	0.5	
Flavour intensity	4.46	3.95	4.17	NS	4.23	4.22	4.13	NS	NS	0.2	
Chewings (number)	24.24°	27.50 ^b	27.19^{b}	*	25.52a	24.37ª	29.04 ^b	NS **	NS	0.2	
Overall	4.62^{a}	3.58^{b}	4.42 ^a	**	4.28a	4.20°	4.15 ^b	*	NS NS	6,2	

Means in the same row not followed by a common letter differ significantly, Sig: Significance; NS (non significance) (p<0.05), ** (p<0.01), *** (p<0.001). MSE: mean square of the error. LD= M. longissimus dorsi. ST= M. semitendinosus, SS= M. supraspinatus; Regions: 1 (cranial in LD, and dorsal in ST and SS), 2 (middle), 3 (candal in LD, and ventral in ST and SS).

Effects of different muscles and regions inside these muscles on textural and sensorial parameters are displayed in Table 2. According to compression measurements by TPA method, each muscle presented a same gradient of hardness are chewiness. The region 3 showed a higher hardness (p<0.01) and chewiness (p<0.05), although only for SS and ST muscles (ventral region), while the LD muscle did not vary within the region in this study. These differences corresponded with the most number of chewings until swallowing (p<0.01) and a less overall (p<0.05) by sensorial analysis. There were no differences in sensorial hardness between the three muscles studied, similar to that found by Searls et al., (2005) who did not observe tenderness differences among steak locations in four beef muscles.

The results based on sensorial analysis showed a significant difference between the muscles studied. LD was more tender (p<0.01) and with less springiness (p<0.05) than ST and SS muscles. Also LD showed less chewiness by IPA (p<0.001) than ST and SS muscles. Low compression rates, in raw meat, are clearly influenced by agoing in the development of meat tenderness and high compression rates demonstrate the influence of connective tissue composition (Campo et al., 2000). ST muscle displayed a smaller juiciness (p<0.01) and overall (p<0.01) than LD and SS muscles principally due to a smaller percentage of intramuscular fat and a higher content of total collagen. Springiness, fat sensation and flavour intensity were no different between muscles studied.

Conclusion

Cranial region in LD and dorsal in ST and SS showed the most overall. Between muscles we observed that ST was sensory depreciated principally due to its lower fat content, higher total collagen value and greater hardness and chewiness by TPA method than others, which was a good indicator of sensory hardness.

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