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FRAGMENTATION INDEX OF RAW MUSCLE AS A TENDERNESS PREDICTOR OF STEAKS FROM US GOOD AND US STANDARD STEER AND BULLOCK CARCASSES

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Summary

Thirty steer (10 US Good, 20 US Standard) and 10 bullock carcasses (one US Good, nine US Standard) were selected from two commercial meat packing firms and aged for 10 to 14 days in a 2 C cooler. Each carcass was assigned scores for the various USDA quality and yield grade factors during a 48- to 120-hr post-mortem selection period. Steaks containing the *longissimus* muscle were obtained from the anterior end of the short loin and cooked to 70 C. They were then measured for tenderness with the Warner-Bratzler shear and evaluated by a trained eight-member sensory panel. Fragmentation index (FI) was determined on fresh and frozen raw *longissimus* muscle at each of three posthomogenization residue fraction drying periods (10 min, 40 min and 24 hr). Sarcomere length also was determined. Simple correlation coefficients relating FI (10 min, 40 min and 22 hr) to tenderness rating for bullocks were: (1) $-.74$, $-.75$ and $-.72$, respectively, for fresh muscle and (2) $-.70$, $-.66$ and $-.69$, respectively, for frozen muscle FI. Simultaneous consideration of carcass physical traits, sarcomere length and eight FI values accounted for 78.7 and 72.6% of the observed variation in shear force value and tenderness rating of steaks from the 30 US Good and US Standard steer carcasses. FI determined from fresh *longissimus* muscle accounted for 48.4 and 44.5% more of the variation in cooked meat tenderness than did carcass physical traits and sarcomere length. The best prediction equations for fragmentation of either fresh or

frozen raw muscle accounted for over 53% of the observed variation in cooked meat shear force values; this degree of precision can be achieved in approximately 15 min of laboratory time. These data indicate that FI of raw muscle can be used for stratification of low grading bullock and steer carcasses according to tenderness level of cooked loin steaks.

(Key Words: Fragmentation, Tenderness, Beef, US Good, US Standard, Bullock.)

Introduction

Recently, much effort has been extended to clarify the relationship between raw muscle fragmentation and cooked meat tenderness (Moller *et al.*, 1973; Berry *et al.*, 1974; Reagan *et al.*, 1975; Olson and Parrish, 1977; Culler *et al.*, 1978; Davis *et al.*, 1979; Calkins *et al.*, 1980). A variety of procedures and quantitation methods exist for fragmenting raw muscle: Davis *et al.* (1980) expressed a fragmentation index (FI) based on the residue weight of fragments more than 250 μm in size; Berry *et al.* (1974) reported the percentage of short fiber fragments observed microscopically, and Moller *et al.* (1973) and Olson *et al.* (1976) measured absorbance of a myofibril suspension.

Commercial application of fragmentation would require the use of a rapid simple and accurate procedure. Calkins *et al.* (1980) reported success in identifying tender carcasses within the US Commercial or US Utility grades with the FI procedure developed by Davis *et al.* (1980). The researchers concluded that fragmentation of frozen, raw *longissimus* muscle accounted for 56.6% of the observed variation in shear force value of cooked loin steaks.

Since there is considerable variation in the palatability of bullock carcasses (USDA, 1972) and since the US Standard grade has been

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broadened (USDA, 1976), an economic advantage might be realized if certain tender carcasses within these classifications could be objectively identified. The objective of the present study was to determine the relationship between fragmentation of raw muscle (fresh and frozen) and cooked meat tenderness of steaks from bullock and low quality steer carcasses.

Experimental Procedure

Thirty steer (10 US Good, 20 US Standard) and 10 bullock carcasses (nine US Good, one US Standard) were evaluated for grade factors at 48- to 120-hr postmortem and transported

to the University of Tennessee meat laboratory. After an aging period of 10 to 14 days in a 2 C cooler, three steaks were removed from the anterior end of the short loin. Steak A (.65 cm thick) was obtained for fragmentation and sarcomere length determination. Palatability and shear force value were determined for steaks B and C (3.2 cm thick).

FI for fresh and frozen raw muscle was determined by the procedure of Davis *et al.*, (1980). Dorsal blade depth was maintained at 1 mm (Calkins and Davis, 1978) below the surface of the solution, and weights of residue fractions were taken at 10 min, 40 min (22 C

TABLE 1. MEANS AND COEFFICIENTS OF VARIATION FOR PHYSICAL TRAITS, PALATABILITY ATTRIBUTES, SHEAR FORCE VALUE AND FRAGMENTATION MEASURES FOR BULLOCK AND STEER CARCASSES

Trait	Bullocks ^a		Steers ^a	
	Mean	CV ^b	Mean	CV ^b
Physical traits				
Marbling ^c	21.2	33.78	27.0	19.07
Lean color ^d	5.2	34.88	6.1	13.64
Lean firmness ^d	3.4	60.75	5.3	25.88
Lean texture ^d	5.9	44.09	7.1	16.55
Fat thickness, mm	1.4	131.74	6.4	50.25
Carcass weight, kg	227.6	16.18	205.0	13.13
Sarcomere length, μ m	1.76	8.93	1.81	7.33
Palatability attributes^e				
Tenderness rating	4.5	27.70	5.6	22.17
Juiciness rating	4.9	11.62	4.8	11.21
Connective tissue rating	6.8	12.01	7.6	4.30
Flavor desirability rating	4.5	27.42	5.8	11.10
Overall satisfaction rating	4.4	26.51	5.3	17.41
Shear force value, kg	5.39	40.71	3.97	42.36
Fragmentation index^f				
Fresh, 10 min	235.4	55.13	147.3	65.04
Fresh, 40 min	205.1	55.43	126.8	67.27
Fresh, 22 hrs	48.6	53.47	32.8	64.29
Fresh, filtrate volume, ml	54.3	5.49	55.3	4.20
Frozen, 10 min	346.3	50.81	271.3	48.65
Frozen, 40 min	301.7	49.12	235.6	49.14
Frozen, 22 hr	75.1	51.98	64.1	52.62
Frozen, filtrate volume, ml	52.0	5.31	53.3	4.54

^aBullocks: one US Good, nine US Standard; steers: 10 US Good, 20 US Standard.

^bCV = coefficient of variation.

^cMean based on 100-unit scale (70 = Slightly abundant^{oo}, 30 = Slight^{oo}).

^dMeans based on eight-point scoring scale (8 = light grayish red, very firm or very fine texture; 4 = moderately dark red, slightly soft or slightly coarse).

^eMeans based on eight-point rating scale (9 = extremely tender, extremely juicy, extremely desirable flavor, no connective tissue or extremely desirable overall; 1 = extremely tough, extremely dry, extremely undesirable flavor, abundant amount of connective tissue or extremely undesirable overall).

^fFragmentation index = 100 \times weight (grams) after air drying at 22 C (10 min and 40 min) and oven drying at 35 C (22 hr).

air dry) and 22 hr (35 C oven dry) for both fresh and frozen raw *longissimus* muscle. Filtrate volume was also recorded.

Samples for sarcomere length were blended in an Osterizer Cycle-Blend for 90 seconds. The suspension medium was 4% formalin (25 ml). A light microscope ($\times 1,500$) and filar micrometer were used to measure 10 sarcomeres on each of 12 myofibrils.

Sensory evaluation (steak B) and shear force value (steak C) were obtained on steaks cooked to an internal temperature of 70 C on individual, preheated broiling units. Final internal temperature was monitored by thermocouples. Steaks B and C, previously frozen at -31 C and stored at -18 C, were thawed for 24 hr in a 5 to 7 C cooler before being cooked. A trained, eight-member sensory panel evaluated tenderness, juiciness, connective tissue amount, flavor desirability and overall satisfaction for each sample, using eight-point rating scales (8 = extremely tender, extremely juicy, no connective tissue, extremely desirable flavor or extremely desirable overall; 1 = extremely tough, extremely dry, abundant connective tissue, extremely undesirable flavor or extremely

undesirable overall). Steak C was cooled to 25 C, and four 1.3-cm cores were removed and sheared in duplicate on a Warner-Bratzler shear.

Data were analyzed according to guidelines outlined by Steel and Torrie (1960).

Results and Discussion

Mean values for various physical, palatability, shear and fragmentation measures on the steer and bullock carcass samples are presented in table 1. The 10 US Good and 20 US Standard steer carcasses were pooled into one group of 30, because an analysis of variance for USDA quality grade with sensory tenderness and shear force value as dependent variables resulted in nonsignificant F values of .00 and .54. Among steaks from the low grading steer carcasses ($n = 30$), increased tenderness was associated with decreased FI values for fresh and frozen raw muscle at all drying times (table 1). Comparison of data for steaks from steer and bullock carcasses shows the samples from steer carcasses (1) were rated 1.1 tenderness units higher by the sensory panel, (2) were 1.42 kg lower in shear force value and (3) had an FI 11.0 to 88.1

TABLE 2. SIMPLE CORRELATION COEFFICIENTS RELATING FRAGMENTATION MEASURES TO PHYSICAL TRAITS, PALATABILITY ATTRIBUTES AND SHEAR FORCE VALUE FOR BULLOCK CARCASSES^a

Trait	Fragmentation of fresh muscle ^b				Fragmentation of frozen muscle ^b			
	10 min	40 min	22 hr	Filtrate volume	10 min	40 min	22 hr	Filtrate volume
Physical traits								
Marbling	.16	.15	.20	-.26	-.12	-.11	-.09	.20
Lean color	-.65	-.67	-.59	.72	-.55	-.51	.37	-.68*
Lean firmness	.21	.20	.27	.22	-.10	-.07	-.04	.17
Lean texture	-.77**	-.78**	-.75*	.61*	-.51	-.45	-.41	.68*
Fat thickness	-.23	-.24	-.19	.24	-.11	-.06	.02	.31
Carcass Weight	.47	.45	.51	-.50	.54	.55	.61	-.38
Sarcomere length	-.42	-.44	-.37	.33	-.37	-.30	-.23	.59
Palatability attributes								
Tenderness rating	-.74*	-.75*	-.72*	.52	-.70*	-.66*	-.69*	.73*
Overall satisfaction rating	-.73*	-.74*	-.72*	.51	-.65*	-.61	-.63	.73*
Shear force value	.38	.39	.36	-.18	.53	.50	.52	-.56

^aN = 10 (one US Good, nine US Standard).

^bFragmentation index = $100 \times$ weight (grams) after air drying at 22 C (10 min and 40 min) and oven drying at 35 C (22hr).

* P < .05.

**P < .01.

units higher. These data indicate that the FI can segment slightly tender (steer) from slightly tough (bullock) cooked steaks with the use of fresh or frozen raw muscle.

FI of raw muscle from bullock carcasses was significantly related to tenderness rating in seven of eight comparisons (table 2). Subjective lean color and lean firmness scores for bullock carcasses (one US Good, nine US Standard) were significantly related ($P < .05$) to two to five measures of fragmentation (table 2). These data indicate that a bullock carcass with a youthful-colored, fine-textured lean will most likely be rated tender and possess a lower FI than a bullock carcass with dark, coarse-textured lean. The significant association between FI and lean texture (table 2) is in agreement with values reported by Calkins *et al.* (1980) for US Commercial and US Utility beef.

Simple correlation coefficients relating fragmentation measures to physical traits and sensory ratings of the US Good and US Standard steer carcasses are presented in table 3. All fragmentation values (fresh and frozen muscle) were significantly correlated with the sensory tenderness rating and shear force value (table 3). The correlation coefficients in tables 2 and 3 for lower grading, A maturity carcasses are in

agreement with the coefficients between FI and tenderness reported by Calkins *et al.* (1980) for US Commercial and US Utility beef and by Davis *et al.* (1980) for US Choice, US Good and US Commercial carcasses. The coefficients also agree with values between MFI and tenderness reported by Olson and Parrish (1977). Culler *et al.* (1978) and Parrish *et al.* (1979).

Twelve physical and histological variables accounted for approximately 75% of the observed variation in shear force value or tenderness rating of steaks from the 30 US Good and US Standard steer carcasses (table 4). Physical traits (marbling degree, lean color, lean texture, carcass weight and fat thickness) and sarcomere length were included first in the model. Results of the analysis (table 4) indicate that physical traits and sarcomere length accounted for approximately 27.5 and 24.3% of the variation in shear force value and tenderness rating. Because of the time required to obtain an FI from a residue fraction oven dried for 22 hr, the following fragmentation values were included next: 10 min FI, 40 min FI, and filtrate volume (fresh and frozen raw muscle). The data (table 4) reveal that fresh muscle fragmentation values explained 48.40 and

TABLE 3. SIMPLE CORRELATION COEFFICIENTS RELATING FRAGMENTATION MEASURES TO PHYSICAL TRAITS, PALATABILITY ATTRIBUTES AND SHEAR FORCE VALUE FOR US GOOD AND US STANDARD STEER CARCASSES (n=30)

Trait	Fragmentation of fresh muscle ^a				Fragmentation of frozen muscle ^a			
	10 min	40 min	22 hr	Filtrate volume	10 min	40 min	22 hr	Filtrate volume
Physical traits								
Marbling	-.13	-.15	-.11	.04	-.19	-.21	-.08	.33
Lean color	-.02	-.04	-.04	.18	.01	.00	-.03	.05
Lean firmness	-.33	-.34	-.31	.42*	-.26	-.25	-.21	.35
Lean texture	-.43*	-.45**	-.43*	.53**	-.29	-.30	-.25	.49**
Fat thickness	-.48**	-.49**	-.47**	.36	-.54**	-.51**	-.48**	.52**
Sarcomere length	.10	.10	.10	-.13	.02	.02	.09	.00
Palatability attributes								
Tenderness rating	-.71**	-.71**	-.70**	.56**	-.66**	-.64**	-.60**	.65**
Overall satisfaction rating	-.68**	-.68**	-.66**	.51**	-.63**	-.62**	-.56**	.64**
Shear force value	.64**	.64**	.63**	-.41*	.73**	.72**	.66**	-.67**

^aFragmentation index = $100 \times$ weight (grams) after air drying at 22 C (10 min and 40 min) and oven drying at 35 C (22 hr).

* $P < .05$.

** $P < .01$.

TABLE 4. COEFFICIENTS OF DETERMINATION AND PARTIAL COEFFICIENTS OF DETERMINATION FOR PREDICTION OF SHEAR FORCE VALUE AND TENDERNESS RATING OF STEAKS FROM US GOOD AND US STANDARD STEER CARCASSES BY CERTAIN PHYSICAL TRAITS AND FRAGMENTATION MEASURES

Model	Shear force value $R^2 \times 100$	Tenderness rating $R^2 \times 100$
Physical traits ^a , fresh muscle fragmentation ^b , frozen muscle fragmentation ^b	78.70	72.60
Physical traits	27.47	24.34
Fresh muscle fragmentation/physical traits ^c	48.40	44.48
Frozen muscle fragmentation/physical traits, fresh muscle fragmentation ^d	2.83	3.78

^aPhysical traits include marbling degree, lean color, lean texture, carcass weight, fat thickness and sarcomere length.

^bFragmentation measures include 10-min index, 40-min index and filtrate volume for fresh and frozen *longissimus* muscle.

^cPartial coefficients of determination accounting for variation in Warner-Bratzler shear force value and tenderness rating which was not previously explained by physical traits.

^dPartial coefficients of determination accounting for variation in Warner-Bratzler shear force value and tenderness rating which was not previously explained by fresh muscle fragmentation and physical traits.

44.48% of the variation in WBS force value and tenderness rating above that variation accounted for by physical traits and sarcomere length.

Regression equations for predicting shear force value of steaks from US Good and US Standard carcasses are presented in table 5. These data were developed in an attempt to identify the percentage of the observed variability in tenderness that could be accounted for by use of one or two measures of fragmentation. The best single fragmentation measure for use of fresh and frozen raw muscle accounted for 41.56 and 53.09% of the observed variation in shear force value. An additional 16.45% (fresh muscle) and .08% (frozen muscle) precision was attained when a second fragmentation variable was used to predict shear force value (table 5). With precision and laboratory time considered, the best prediction equations for fresh (10 min FI and filtrate volume) and frozen (10 min FI) raw muscle accounted for over 53% of the observed variation in cooked meat shear force values. This degree of precision can be achieved in approximately 15 min (including homogenization, filtration and fragment drying time periods). To facilitate the routine determination of FI, it may be more desirable to use fresh rather than frozen raw muscle, especially since the 5-min thawing time

period can be eliminated and, as evidence in table 4 indicates, fragmentation of frozen muscle adds less than 4% precision to prediction of cooked meat tenderness. Conversely, in a study of US Commercial and US Utility carcasses, Calkins *et al.* (1980) has reported that

TABLE 5. REGRESSION EQUATIONS FOR PREDICTION OF SHEAR FORCE VALUE OF STEAKS FROM US GOOD AND US STANDARD STEER CARCASSES WITH USE OF CERTAIN FRAGMENTATION MEASURES

State of raw muscle	Variables ^a	$R^2 \times 100$	SEE ^b
Fresh	1, 3	58.01	2.49
Fresh	1	41.56	2.88
Frozen	1, 2	53.17	2.62
Frozen	1	53.09	2.58

^aVariable code for fragmentation measures:

1 = FI after air drying of residue fraction at 22 C for 10 minutes.

2 = FI after air drying of residue fraction at 22 C for 40 minutes.

3 = Filtrate volume.

^bSEE = standard error of the estimate.

FI determined from frozen *longissimus* muscle accounted for 18.6 to 23.8% more of the observed variation in cooked meat tenderness than did FI of fresh muscle.

On the basis of (1) the data presented herein for lower grading, A maturity steer and bullock carcasses; (2) results obtained by Calkins *et al.* (1980) for C maturity carcasses, and (3) data reported by Stiffler and Ray (1979) for short scrotum and steer carcasses, it appears that the FI procedure (Davis *et al.*, 1980) may have potential for industrial application and (or) as a routine meat science research technique. The above uses are suggested since fragmentation can be performed in approximately 15 min on raw muscle representing a wide range of carcasses and account for over 53% of the observed variation in cooked meat tenderness.

Conclusions of our major findings in the present study were that: (1) FI of raw muscle for low grading bullock carcasses was significantly related to tenderness rating of cooked loin steaks. (2) Fragmentation values were superior to physical traits and sarcomere length in segmenting low grading steer carcasses according to tenderness rating or shear force value of cooked loin steaks. (3) Neither FI of raw muscle nor sensory tenderness and shear force value of cooked steaks were successful in segmenting US Good from US Standard steer carcasses. (4) The best prediction equations for fragmentation of either fresh or frozen raw muscle accounted for over 53% of the observed variation in cooked meat shear force values, a degree of precision which can be achieved in approximately 15 min of laboratory time.

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