

1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29

Running head: Correlation of loin firmness with pork quality

Correlation of fresh muscle firmness with sensory characteristics of pork loins destined for a quality focused market¹

E.K. Arkfeld*, S. Mancini**†, B. Fields‡, A.C. Dilger*, D.D. Boler*²

*Department of Animal Sciences, University of Illinois, Urbana-Champaign, IL 61801

†Department of Veterinary Sciences, University of Pisa, Pisa, Italy 56124

‡PIC, Hendersonville, TN 37075

¹Acknowledgements: Gratitude is expressed to M. A. Tavárez, B. M. Bohrer, M. F. Overholt, S. E. Curtis, and R. Schmitt for technical assistance in the completion of this research.

²Corresponding author: dboler2@illinois.edu (D.D. Boler)

30 **ABSTRACT:** Production of pork for quality driven export markets offers economic incentive.
31 Pork processors use subjective firmness as a sorting tool for loins intended for high-quality
32 export. The objectives of this study were to determine: 1) durometer efficacy in muscle, 2) if
33 firmness on one portion of the loin is indicative of other locations, 3) if 1 d firmness is related to
34 export quality traits, and 4) if variation in firmness is explained by mechanistic measures.
35 Subjective firmness scores (1 = extremely soft; 5 = extremely firm) were determined by a trained
36 individual d 1 (initial time point) postmortem. Loins (NAMP #414 Canadian back, N=154) were
37 **wet** aged for 28 d at 1.7°C. On d 28, a panel of 4 individuals assigned firmness scores on the
38 ventral side of the loin at the area of the 10th rib, the anterior half, and the posterior half of the
39 loin. Durometer readings were collected at the area of the 10th rib on the dorsal and ventral side
40 of the loin. Spearman correlation coefficients were computed in SAS (v. 9.3) to account for non-
41 normality of categorical data. Subjective firmness measures at d 28 at the 10th rib and on the
42 anterior portion of the loin did not correlate ($P \geq 0.21$) with whole loin durometer readings on the
43 dorsal or ventral portion of the loin, nor the average of the whole loin values. Subjective firmness
44 (d 28) at the 10th rib accounted for 38.44% ($r = 0.620$) and 48.30% ($r = 0.695$) of the variation in
45 firmness at the anterior portion of the loin and the posterior portion of the loin, respectively ($P \leq$
46 0.05). One d subjective firmness measures correlated with 28 d WBSF measures ($r = 0.174$; $P =$
47 0.03), but did not significantly correlated with sensory characteristics ($P \geq 0.08$). Purge tended
48 to correlated with 1 d firmness ($r = 0.136$; $P = 0.10$), however drip and cooking loss, 24 h and 28
49 d pH, and soluble and insoluble collagen content were not correlated ($P \geq 0.34$). Firmness
50 measures collected in the production facility (d 1) negatively correlated with IV ($r = -0.199$; $P =$
51 0.02) yet no 28 d subject firmness measures were correlated with IV ($P \geq 0.33$). When loins not
52 achieving export standards are removed from the population, 1 d firmness was not correlated to

53 export quality or sensory characteristics (d 28). Differences in firmness were not explained by
54 mechanistic measures. Inconsistencies among subjective and objective firmness measures
55 suggest the durometer may not be an appropriate way to determine firmness.

56

57

58

59

60

61

62

63

64

65

66

67

68

69

70

71

72

73

74

75

76

77 **Key words:** export, firmness, loin, pork, quality, sensory

78

INTRODUCTION

79

80

81

82

83

84

In 2014, Mexico imported the greatest amount of us pork (680,842 metric tons; \$1.558 billion), while Japan was the largest importer of U.S. pork on a value basis (468,561 metric tons; \$1.932 billion; Masker, 2015). Japanese meat processors rank pork eating quality second only to food safety (Murphy et al., 2015). Therefore, one challenge for U.S. pork processors seeking to export to high-quality markets like Japan is to determine eating quality of the product prior to shipping without harming the integrity of whole muscle products.

85

86

87

88

89

90

91

92

93

94

95

96

A variety of mechanical methods are available to predict quality (pH meter, colorimeter, etc.), but these methods are not a viable option at production speed (> 1000 pigs/h) in a U.S. production facility. Consequently, many quality decisions for fresh loins are based on assessments of subjective color, marbling, and firmness by facility personnel. Studies have evaluated the correlations of subjective firmness to other pork quality traits (Huff-Lonergan et al., 2002; Boler et al., 2010). However, these studies have not focused on subjective firmness as a determining factor for palatability of loins destined for export. Others have evaluated mechanical firmness of muscle (Rincker et al., 2007) and adipose tissue (Seman et al., 2013). Seman et al. (2013) demonstrated that durometers are effective at evaluating tissue firmness in pork bellies, but less is known about the durometers efficacy in muscle tissue. As such, little information exists regarding appropriateness of using loin firmness as a sorting tool to predict quality of exported loins.

97

98

99

100

Therefore, this study had four primary objectives: 1) to determine if durometer measures correlate with subjective firmness scores, and thus have potential efficacy in a commercial facility, 2) to determine if firmness at various anatomical locations throughout the loin is an accurate predictor of firmness in other loin locations, 3) to determine if firmness 1d postmortem

101 is related to export quality traits, and 4) to determine if variation in firmness can be attributed to
102 variability in mechanistic measures. Although correlations have been made between firmness
103 and other quality traits, we hypothesize that by removing the lowest quality loins from a
104 population of loins, variation in firmness will be reduced. Thus, in high-quality export loins,
105 firmness is not an accurate indicator of pork quality and may not be the most appropriate way to
106 select pork loins for export.

107 MATERIAL AND METHODS

108 Postmortem samples were obtained from a federally inspected slaughter facility therefore
109 no Institutional Animal Care and Use Committee approval was necessary.

110 *Processing Facility Data Collection*

111 Loins (N = 154) selected for this experiment were derived from PIC337-sired pigs (PIC,
112 Pig Improvement Company, Hendersonville, TN). Hot carcass weight, backfat depth, and loin
113 depth were measured after slaughter of the pigs using a Fat-O-Meater system (Fat-O-Meater
114 measurements, SFK Technology Fat-O-Meater, Herlev, Denmark). Estimated percent lean was
115 calculated using a proprietary facility equation. Carcasses were fabricated into primal cuts at
116 approximately 24 h postmortem. Boneless loins were evaluated for subjective quality measures
117 online 1 d postmortem on the ventral surface of the loin after the backribs were removed. Color
118 was evaluated online approximately 3-5 min after backribs were removed using a 6 point scale
119 (Japanese color scale), marbling was evaluated using a 10 point scale (NPPC, 1999; 1 = 1.0%
120 intramuscular lipid, 10 = 10.0% intramuscular lipid), and firmness was evaluated using a 5 point
121 scale (NPPC, 1991; 1=soft; 5=very firm) through standard facility procedures of folding the
122 whole loin. Objective L*, a*, and b* measurements were collected using a Minolta CR400
123 Colorimeter (C light source, 2° observer, 10 mm aperture). Ultimate pH was recorded using a

124 MPI pH meter with glass tipped electrode (Meat Probes Inc., Topeka, KS), the probe was
125 inserted at approximately the 10th rib location on the boneless loin. Iodine value of jowl and
126 belly adipose tissue was measured using a Bruker NIR (Billerica, MA). Loins were individually
127 packed in vacuum-packaging and shipped under refrigeration to the University of Illinois Meat
128 Science Laboratory.

129 *Aged Loin Evaluation*

130 Loins arrived at the University of Illinois Meat Science Laboratory under refrigeration at
131 1.7 °C. Loins were aged, **in vacuum-sealed packages**, until 28 d postmortem at 4 °C to account
132 for an estimated time it would take loins to arrive at their final export destination. Aged loins
133 were weighed in their package and weighed after removal from their package. Dried package
134 weight was determined as the average of a random selection of 10% (16 packages) of the
135 vacuum-packaged bags used in the study and subtracted from the packaged loin weight. Purge
136 loss was calculated as weight lost in purge as a percentage of packaged weight.

137 *Fresh Loin Firmness Measurements*

138 Loin flop distance was determined by placing the loin, ventral side down, on a bar and
139 measuring the distance between the inside edges of both ends of the loin. A trained panel (n=4)
140 assigned firmness scores to whole loins on a scale of 1 to 5; with one being very soft and 5 being
141 very firm (NPPC, 1991). Aged (28 d) fresh loin firmness was evaluated at several location of the
142 loin and on multiple, individual chops to gain an understanding of the variation on firmness
143 throughout the loin. Subjective firmness of the ventral side of the loin was assessed at three
144 locations: mid-point (at approximately the 10th rib), anterior (half the distance between the 10th
145 rib and anterior end of the loin), and posterior (half the distance between the 10th rib and

146 posterior end of the loin). Durometer measurements (objective firmness with greater numbers
147 indicating a firmer product; DD-100-000-S with removable stainless steel barrel; Check-Line,
148 Cedarhurst, NY) were determined at approximately the 10th rib on both the dorsal and ventral
149 sides of the loin with an approximate temperature of 4 °C. Durometer measures on the ventral
150 portion of the loin were evaluated on fresh muscle tissue and measures on the dorsal portion of
151 the loin were evaluated on loins trimmed to the epimysium. Dorsal and ventral durometer
152 measures were averaged for a whole loin durometer firmness score. Loins were trimmed to
153 expose a fresh cut surface at approximately the 10th rib, where the trained panel evaluated
154 firmness on the cut surface. Chops (2.54 cm thick) were cut assigned to assays (Warner-Bratzler
155 shear force and sensory evaluation) in a consistent order to minimize variation due to loin
156 location among loins. Subjective firmness and durometer readings were collected on lean tissue
157 of these chops and then, they were held frozen at -4° C in a vacuum package bag for further
158 analyses. Additionally, 1.25 cm thick chops were collected for drip loss and 7.6 cm section was
159 collected for intramuscular fat iodine values.

160 *Color, Marbling, and pH Measurements*

161 Objective and subjective color readings were collected on the cut surface of the loin
162 immediately after facing the loin (pre-bloom) at approximately the 10th rib. Objective L*, a*,
163 and b* measurements were collected using a Konica Minolta CR-400 colorimeter (Minolta
164 Camera Company, Osaka, Japan; D65 light source, 0° observer, 8 mm aperture). Subjective
165 color scores were assigned on a scale of 1 to 6 (NPPC, 1999), where 1 represented a pale pinkish-
166 grey color and 6 a dark purplish-red color. Loins were then allowed to bloom (to allow for the
167 conversion of deoxymyoglobin to oxymyoglobin) for at least 20 min, and color was measured
168 again using the same protocol as above. Subjective marbling scores were assigned to loins after

169 the 20 min bloom time using a scale of 1 to 10 (NPPC, 1999), where 1 = 1.0% intramuscular
170 lipid and 10 = 10.0% intramuscular lipid. Ultimate pH was measured using a MPI pH meter
171 (Meat Probes Inc., Topeka, KS; 2 point calibration at pH 4 and 7). The pH probe was inserted
172 on the cut surface towards the posterior end of the loin.

173 *Drip Loss*

174 Chops cut 1.25 cm thick were used for determination of drip loss. An initial weight was
175 recorded, and chops were suspended in Whirl Pak bags for 24 h at 4 °C. Final weight was
176 collected after 24 h and drip loss was calculated as: $[(\text{initial weight} - \text{final weight}) / \text{initial}$
177 $\text{weight}] \times 100$.

178 *Warner-Bratzler Shear Force (WBSF) and Cook Loss*

179 Samples for WBSF were thawed at 4° C for 24 h. Chops were trimmed of excess
180 subcutaneous fat, weighed, and cooked to 70°C on Faberware open hearth grills (Model 455 N,
181 Walter Kiddie, Bronx, NY. Internal temperatures were monitored using thermocouples (Type T,
182 Omega Engineering, Stamford, CT) connected to a digital scanning thermometer (Model 92000-
183 00 Barnant Co., Barington, IL). Chops were weighed after tempering to approximately room
184 temperature and cook loss was calculated as: $[(\text{raw weight} - \text{cooked weight}) / \text{raw weight}] \times 100$.
185 After cooling to approximately 22 °C, six 1.25 cm cores were removed parallel to the orientation
186 of muscle fibers and sheared using a Texture Analyzer TA.HD Plus (Texture Technologies
187 Corp., Scarsdale, NY/Stable Microsystems, Godalming, UK) with a blade speed of 3.3 mm/s and
188 100 kg load cell capacity. The average of the 6 cores were reported as WBSF values.

189 *Sensory Panel*

190 **Trained** panelists (n=6) evaluated samples for tenderness, chewiness, juiciness, and off
191 flavor. Chops were trimmed of subcutaneous fat and cooked in the same manner as chops for
192 WBSF. No greater than 8 samples were served per panel; allotment of chops to panel was
193 random. Two cubes (1 cm x 1 cm x 2.54 cm) were served to each panelist under red lighting. A
194 15 cm anchored scale was used with a low degree of each trait on the left side of the scale (0)
195 and a high degree of each trait on the right side (15; Smith et al., 2011; Arkfeld et al., 2015).

196 *Proximate Composition*

197 Chops for proximate composition were thawed, trimmed of subcutaneous fat, and
198 homogenized in a food processor (Cuisinart, East Windsor, NJ). Methods were followed as
199 described by Novakofski et al. (1989). Briefly, moisture and extractible lipid analyses were
200 performed in duplicate. Samples were dried at 110° C for at least 24 h and extracted in an
201 azeotropic mixture of warm chloroform:methanol. Protein concentrations were determined by
202 measuring N content using the combustion method (Association of Official Analytical Chemists,
203 2000; model TruMac, method 990.03, LECO Corp., St. Joseph, MI).

204 *Soluble and Insoluble Collagen Content*

205 Chops for collagen determination were trimmed of external fat, frozen in liquid nitrogen,
206 and then ground to a powder using a blender (Waring Commercial Blender Model HGB2WT53,
207 Stamford, CT). Soluble and insoluble collagen content procedures were adapted from protocol
208 outlined by Hill (1966). Duplicate 3.0 gram samples were weighed into 50 mL polyethylene
209 tubes, 16 mL of ¼ strength Ringer's solution was added. Samples were placed in a 77 °C water
210 bath for 70 min, with samples shaken every 10 min. Next, samples were centrifuged at 5200 x g
211 for 10 min and the soluble fraction was decanted into a flask labeled soluble fraction through

212 filter paper (Qualitative P8, 15.0 cm diameter, Fisher Scientific, Pittsburgh, PA). Eight mL of $\frac{1}{4}$
213 strength Ringer's solution was added to the remaining sample in the polyethylene tube and
214 sample was centrifuged and decanted in the same manner. Sample remaining in the tube
215 (insoluble fraction) was removed using a metal spatula into a flask labeled as insoluble fraction.
216 One-half of an extra low lint task wipe (Kimwipe EX-L, Kimberly-Clark; Dallas, TX) was used
217 to remove any remaining insoluble fraction from the polyethylene tube. The Kimwipe and filter
218 paper used to decant the soluble fraction were placed in the insoluble flask for their respective
219 sample. Twenty-five mL of 6 N HCl were added to each flask and the flasks were placed in a
220 110 °C oven for at least 12 h. After 12 h 1.0 gram of charcoal was added to the flask, flask was
221 shaken, and then contents were filtered through filter paper (Whatman 2, 150 mm, Sigma-
222 Aldrich, St. Louis, MO). The pH of each sample was buffered to 6.0 ± 0.1 and samples were
223 volumized through Qualitative P8 filter paper. One mL of each sample was combined with 2 mL
224 of isopropanol and vortexed. One mL of oxidant solution (1 volume chloramine T to 4 volumes
225 of acetate citrate buffer) was added to each sample and vortexed. Exactly 4 min later, 4 mL of
226 Elrich's solution (15.8% dimethylaminobenzoaldehyde·HCl₄ and 84.2% isopropanol) was added
227 to the sample and vortexed. Samples were placed in a 60 °C water bath for 25 min, and then a
228 cool water bath for 5 min. Two hundred μ l of each tube was plated along with **hydroxyproline**
229 standards of 0, 4, 8, 12, 16, 20, 24, 28, 32, and 36 μ g/mL prepared in the same manner as
230 samples. Plates were read at an absorbance of 558 nm, and collagen content was determined as
231 $[(\mu\text{g/mL}) (\text{dilution factor}) (\text{constant})] / [\text{sample weight} \times 1000]$, with constants of 7.52 and 7.25
232 for soluble and insoluble samples, respectively.

233 *Intramuscular Fat Iodine Value*

234 Intramuscular lipid was extracted from muscle tissue using the procedure of Folch et al.
235 (1957). Fatty acid methyl esters (FAME) were converted from lipid using the AOAC official
236 method C3 2-66 (1998). The resulting FAME were analyzed using a gas chromatograph
237 (Hewlett Packard 5890 series II) equipped with an auto-sampler and a DB-wax capillary
238 column (30m x 0.25mm x 0.25 μ m film coating; Agilent Technologies, Santa Clara, CA). The
239 equipment was operated under constant pressure (1.30 kg/cm²) using helium as the carrier gas
240 and a 100:1 split ratio. Temperature of the injector was held constant at 250 °C and temperature
241 of the flame-ionization detector was held at 260 °C. The oven was operated at 170 °C for 2 min
242 and then increased 2 °C per min up to 240 °C when this temperature was maintained for 8 min.
243 Chromatographs were integrated using Agilent Chemstation Software for gas Chromatographs
244 systems (Version B.01.02, Agilent Technologies, Inc.). Peaks were identified using a gas
245 chromatograph reference standard (GLC 461 A, Nu-check-prep, Elysian, MN). Fatty acids were
246 normalized such that the area of each peak was represented as the percentage of the total area.
247 Iodine values were calculated using fatty acid profile data with the following AOCS (1998)
248 equation: IV = C16:1 (0.95) + C18:1 (0.86) + C18:2 (1.732) + C18:3 (2.616) + C20:1 (0.785) +
249 C22:1 (0.785).

250 *Statistical Analysis*

251 Data were analyzed using the correlation procedure of SAS (v. 9.3, SAS Institute Inc.,
252 Cary, NC). **There were 154 replications in this study.** Spearman correlation coefficients were
253 used to account for the non-normality of categorical firmness data. Percent variation accounted
254 for by firmness scores was calculated as the correlation coefficient (r), raised to the second
255 power and multiplied by 100 (r² * 100). **Relationships were considered statistically significant at**
256 **the P \leq 0.05 level, and trending at the P > 0.05 to P \leq 0.10 level.**

257

RESULTS AND DISCUSSION

258 Loins used in the study were representative loins selected for export in regards to 1 d color
259 (range 2.5-4), marbling (range 1-3), and firmness (range 2-5; Table 1). Although carcass weight
260 of the current population was similar to previous work investigating pork quality correlations by
261 Huff-Lonergan et al. (2002), carcasses from pigs of the current population had approximately 15
262 mm less fat over the 10th rib. Further, subjective color and firmness scores (both 1 d postmortem)
263 were similar in both populations, but the mean marbling score was 1.85 units greater and pH was
264 0.21 units greater in the Berkshire x Yorkshire F1 population used by Huff-Lonergan et al.
265 (2002). Due to differences in sensory scales used between the two projects direct comparison of
266 sensory data cannot be made.

Durometer Efficacy

268 Processing facility decisions of which loins to export to quality driven markets are based
269 on the following quality criteria 1) lean color, 2) firm muscle, and 3) sufficient marbling
270 (Johnson, 2008). While validated objective measures exist to determine both loin color and lipid
271 percentage, no objective measures to determine muscle firmness have been readily adopted by
272 industry. Durometers have been used to determine fat firmness in bellies (Seman et al., 2013),
273 but results of durometer use on muscle have not been reported. Therefore, in order to validate
274 use of the durometer for loin firmness evaluation, durometer measurements were correlated with
275 subjective firmness measurements.

276 Day 1 subjective firmness (determined at the processing facility) did not correlate with
277 durometer measures of the whole aged loin muscle ($P \geq 0.12$; Table 2). Subjective d 1 firmness
278 tended to correlate with durometer readings on chops used for sensory ($P = 0.07$; $r = 0.146$), but

279 did not correlate with chops used for WBSF ($P = 0.49$). Subjective firmness measures at d 28 at
280 the 10th rib and on the anterior portion of the loin did not correlate with whole loin durometer
281 readings on the dorsal or ventral portion of the loin, nor the average of the whole loin values ($P \geq$
282 0.21). There was, however, a tendency for the subjective measures on the posterior portion of
283 the loin to correlate with mid-point (10th rib) dorsal ($P = 0.08$; $r = 0.142$) and ventral ($P = 0.06$; r
284 $= 0.151$) durometer readings which contributed to a significant correlation of subjective firmness
285 of the posterior portion of the loin with average whole loin durometer measures ($P = 0.04$; $r =$
286 0.170).

287 In general, durometer readings from the area of the 10th rib either individually or when
288 averaged are not well-correlated with subjective firmness scores on any portion of the loin (Table
289 2). A similar result was observed by Swatland (1998), when creating an objective firmness
290 measure using vacuum-induced changes in reflectance of pork loins. Those authors attributed
291 the lack of a relationship between reflectance and firmness to either that objective or subjective
292 measures accounted for different portions of firmness (i.e. elasticity and viscosity), or there was
293 bias in subjective firmness scores by the evaluator due to inconsistencies in pork color
294 (Swatland, 1998). In the present data set, loins were “normal” in color (NPPC color score range:
295 2-4) and neither pale nor dark as in the previous work. Also, a panel of 4 evaluators was used for
296 subjective firmness measures to further eliminate bias. Therefore, while this study does rule out
297 bias as a potential impact on firmness measures, it does not rule out that different portions of
298 firmness (i.e. elasticity and viscosity) may be measured by objective and subjective firmness
299 measures. In the context of the current subjective firmness rating system, the lack of correlation
300 between durometer readings and subjective firmness measures leads to the conclusion that the
301 durometer may not be an appropriate tool for online firmness estimation of pork loins.

302 A widely used commercial application to assess firmness and quality of pork bellies is
303 by the use of belly flop firmness measures. Similar to bellies, it was expected that decreased flop
304 distances would indicate less firm loins. Initial (1 d) and aged (28 d) whole loin subjective
305 firmness traits were correlated with 28 d loin flop ($P \leq 0.01$; Table 3), but no objective firmness
306 measures were related ($P > 0.35$) to loin flop. Subjective and objective whole loin measures are
307 inconsistent with each other, so, the lack of significant correlations between loin flop and
308 objective measures ($P \geq 0.35$) agrees with previous data regarding correlation of belly flop
309 measurements. Trusell et al. (2011) reported no correlation between belly flop measures skin
310 side up, and negative, but inconsistent results across different regions of the belly when
311 measured skin side down compared with mechanical compression measures.

312 *Variation in Firmness Due to Anatomical Location*

313 Firmness determination protocols are plant specific, and therefore a variety of locations
314 are potentially used when determining firmness of loins in the export selection procedure. It is
315 known that firmness varies throughout the loin. Waylan et al. (1998) reported that chops from
316 more posterior portions of the loin are firmer than chops from the anterior portions of the loin.
317 However, it is not well understood if one location within the loin is an accurate predictor of other
318 locations. Therefore, one objective of this project was to determine if firmness in one location of
319 the loin was indicative of firmness in other locations of the loin. Subjective firmness measures at
320 28 d postmortem were significantly correlated with each other. Specifically, firmness at the 10th
321 rib accounted for 38.44% ($r = 0.620$) and 48.30% ($r = 0.695$) of the variation in firmness at the
322 anterior portion of the loin and the posterior portion of the loin, respectively ($P \leq 0.05$; Table 2).
323 Subjective firmness measures on the anterior portion of the loin accounted for 24.4% ($r = 0.494$)
324 of the variation in subjective firmness measures on the posterior portion of the loin. Similarly,

325 durometer readings on the dorsal portion of the loin accounted for 7.1% ($r = 0.266$) of the
326 variation in durometer readings on the ventral portion of the loin ($P < 0.01$). It can be concluded
327 that firmness in one location does predict firmness in other locations of the loin, however this
328 relationship is moderate, and caution should be exercised when comparing results of studies
329 where firmness may be measured in different locations.

330 *Relationship of Export Selection Criteria with 28 d Quality Measures*

331 *Firmness and Sensory Characteristics.* Of all firmness traits measured in this study, initial
332 (1 d) firmness measures are arguably the most important in commerce. Firmness assessed at the
333 processing facility is used in selecting loins for export to high-quality markets. However, few
334 studies have correlated firmness with loins aged over 21 d. In the current study, initial subjective
335 firmness accounted for 3.0% of the variation in WBSF measures ($P = 0.03$; Table 4) while
336 objective durometer readings on the ventral portion of the loin and average whole loin durometer
337 measures tended to correlated with WBSF measures ($P \leq 0.09$). Warner-Bratzler shear force
338 correlated with subjective firmness measures on the chop used to determine WBSF ($P = 0.01$; $r =$
339 0.211). Average whole loin durometer measures and sensory chop objective firmness measures
340 each accounted for 2.7% of the variation in WBSF measures ($P = 0.04$). Previously, WBSF
341 values were correlated with sensory tenderness: sustained tenderness sensory measures
342 correlated with WBSF measures at $r = -0.60$, while initial tenderness sensory measures correlated
343 with WBSF at $r = -0.61$ (Caine et al., 2003). Therefore, the lack of similar correlations in the
344 current study was surprising. In the present study, WBSF was correlated with sensory tenderness
345 at $r = -0.32$ ($P < 0.0001$; data not presented). There was not a significant correlation between
346 sensory tenderness and subjective initial firmness ($P = 0.79$; $r = -0.022$; Table 4), but a
347 significant correlation was present between sensory tenderness and subjective firmness of the

348 anterior portion of the loin ($P = 0.03$; $r = 0.170$). Whole loin objective firmness measures using
349 the durometer did not correlate to sensory tenderness ($P \geq 0.83$). Chewiness, or sustained
350 tenderness, was not correlated with subjective firmness on the posterior portion of the loin ($P =$
351 0.39), yet, was correlated with firmness measures at the 10th rib ($P = 0.03$; $r = -0.180$) and on the
352 anterior portion of the loin ($P = 0.02$; $r = -0.189$). Objective durometer readings from both
353 whole loin and chop measures did not correlate with sensory chewiness ($P \geq 0.14$). Initial
354 firmness tended to explain 2.0% of the variability in sensory juiciness. No significant correlation
355 was observed for sensory juiciness and firmness at the 10th rib and on the posterior portion of the
356 loin, but firmness measures of the anterior portion of the loin tended to account for 1.7% of the
357 variability in sensory juiciness ($P = 0.10$). No subjective firmness measures on the whole loin or
358 on chops were significantly correlated with sensory off-flavor ($P \leq 0.32$). Average whole loin
359 durometer readings accounted for 5.2% of the variation in sensory off-flavor ($P < 0.01$).

360 Davis et al. (1975) proposed correlations between firmness and sensory characteristics. In
361 that study of 403 loins, subjective firmness measures explained between 9.6 to 18.49% of the
362 variation in sensory juiciness, 1.2% of the variation in flavor, 1.4 to 7.8% of the variation in
363 tenderness and 8.4 to 13.7% of the variation in overall satisfaction (Davis et al., 1975). In
364 general, these correlations are greater than correlations presented in the current study, likely due
365 to the added variation of kill location and differing genetics imposed in the Davis et al. (1975)
366 study. In the present study, some firmness measures did account for small variation in sensory
367 characteristics, but these correlations are inconsistent. Consequently, firmness, measured at
368 either 1d or 28 d postmortem with a variety of techniques, is not an accurate predictor of aged
369 pork loin sensory characteristics.

370 ***Firmness, Subjective Color, and Marbling.*** Color is a large determining factor in sorting
371 loins destined for quality driven export markets. Though color can be measured objectively with
372 various colorimeter systems, for sorting, it is measured subjectively. At 1 d color was not
373 correlated with any whole loin subjective firmness measures ($P \geq 0.19$), but was significantly
374 correlated to durometer readings on the dorsal portion of the loin ($P = 0.03$; Table 5).
375 Traditionally, literature reports pork color after a bloom period, however bloom time does not
376 allow for color measurements at line speed, consequently both pre- and post-bloom color were
377 measured at 28 d in this study. No subjective whole loin ($P \geq 0.38$; Table 4) or objective ($P \geq$
378 0.21) firmness measures correlated with pre-bloom color. Pre-bloom subjective color scores
379 were related to durometer measures for texture and sensory chops ($P \leq 0.02$; $r = 0.206$ and 0.192 ,
380 respectively), and tended to be related to firmness of the raw WBSF chops ($P = 0.07$; $r = 0.192$).
381 In regards to subjective firmness measures, firmness of WBSF chops and texture chops were
382 correlated with pre-bloom color ($P < 0.04$; $r = 0.227$ and 0.164 , respectively). Neither whole loin
383 or chop subjective firmness measures correlated with subjective color after a 20 min bloom time
384 ($P \geq 0.19$). Objective sensory chop measures were weakly, but significantly correlated ($P \leq$
385 0.01) to color after the blooming period but the tendency observed pre-bloom for the WBSF
386 chop was no longer present ($P \geq 0.87$). However, these correlations are not as strong as those
387 reported by Huff-Lonergan et al. (2002), who suggested positive correlations between firmness
388 and color of $r = 0.27$, or Boler et al. (2010), who suggested a positive correlation of $r = 0.41$.

389 The final component in evaluating quality of loins for export is marbling, which can be
390 objectively measured by percent extractable lipid, but in commerce, is estimated by visual
391 appraisal. In this population of loins subjective marbling scores ranged from 1 to 3 and
392 extractable lipid from 0.47% to 4.18%. Correlation between these two measures was significant

393 but weak ($P < 0.01$; $r = 0.233$; data not presented). No significant correlations existed between
394 extractable lipid and whole loin subjective firmness measures ($P \geq 0.11$; Table 3). Durometer
395 readings on the dorsal portion of the loin accounted for 2.7% of the variation in extractable lipid
396 ($P = 0.04$), while durometer readings on the ventral portion of the loin and average whole loin
397 durometer measures were not significantly correlated to extractable lipid ($P \geq 0.38$).

398 Although not the emphasis of this experiment, Spearman correlation coefficients are
399 presented on the two remaining export selection criteria (color and marbling) and sensory
400 characteristics (Table 6). Huff-Lonergan et al. (2002) reported significant positive correlations
401 between color and sensory tenderness, as well as color and flavor ratings. Further, in that study,
402 off-flavor was correlated with color. However, in the present study, no significant correlations
403 were reported between 1 d or 28 d color and tenderness, chewiness, and juiciness ($P \geq 0.18$).
404 Similar to Huff-Lonergan et al. (2002) results, a tendency for a negative correlation was present
405 between 1 d color score and sensory off flavor ($P = 0.07$; $r = -0.148$), however this same result
406 was not present between 28 d color and sensory off flavor ($P = 0.90$). Previously, marbling was
407 correlated with tenderness ($r = 0.21$), flavor ($r = 0.20$) and off flavor (-0.15 ; Huff-Lonergan et al.,
408 2002). However, in the present study, sensory characteristics were not correlated with color at
409 either 1 or 28 d of aging ($P \geq 0.33$). Although ranges are not included in the study by Huff-
410 Lonergan et al. (2002), the standard deviations in the present study were lower for 1 d color,
411 marbling and firmness, indicating less variability in the current study and a potential explanation
412 for the lack of correlations between sensory characteristics, and color, marbling, and firmness.

413 ***Relationship Between Mechanistic Traits and Firmness.*** Mechanistic measures are used as
414 indicators of overall quality, palatability, or further processing characteristics. Ultimate pH (24 h
415 postmortem) correlates with color ($r = 0.50$) and marbling ($r = 0.25$; Boler et al., 2010). Previous

416 work suggested that increased pH would result in a swelling of myofibrils (Huff-Lonergan and
417 Lonergan, 2005), which would ultimately result in a firmer product. However, no subjective or
418 objective firmness correlations were observed with 1 d pH ($P \geq 0.13$; Table 5) and no subjective
419 firmness measures were correlated with aged pH measures ($P \geq 0.17$; Table 4). This contrasts
420 with research by both Boler et al. (2010) and Huff-Lonergan et al. (2002) who reported
421 significant, positive correlations between firmness and pH.

422 Firmness has been demonstrated to be negatively correlated with purge loss of pork loins
423 aged for 21 d (Boler et al., 2010). Initial (1 d) firmness measures tended to correlate with aged
424 loin purge ($P = 0.10$; $r = 0.136$; Table 4) such that loins that were firmer on d1 had increased
425 amounts of purge during storage. As loins age and postmortem proteolysis occurs inherent
426 variation in moisture loss occurs (Melody et al., 2004), and may result in a less firm product. It
427 is therefore unsurprising that whole loin subjective firmness measures from the 10th rib, anterior
428 portion, and posterior portion of the loin accounted for 7.2% ($r = -0.268$), 12.2% (-0.349), and
429 9.2% (-0.304), respectively, of the variation of aged loin purge loss ($P < 0.01$). However, at 28 d
430 postmortem, it is expected that the majority of postmortem proteolysis and consequent free water
431 loss is likely nearing completion. This is reflected in the low population average of drip loss
432 percentage in the current project (0.78 ± 0.23 %; Table 1) and the lack of significant correlations
433 between drip loss and subjective whole muscle firmness ($P \geq 0.10$; Table 4). Similar to other
434 traits in this study, durometer readings on the dorsal and ventral portions of the loin did not
435 account for any significant variation in drip loss or purge ($P \geq 0.11$). Furthermore, loin moisture
436 content was not correlated with any firmness measures ($P \geq 0.18$).

437 Collagen content is known to contribute to the variability in tenderness of pork (Wheeler
438 et al., 2000). However, soluble and insoluble collagen content were poor indicators of firmness,

439 with only subjective WBSF chop firmness being correlated with insoluble collagen content ($P =$
440 0.01 ; $r = 0.220$; Table 4), but no other chop or whole loin measurement of firmness. Excessive
441 unsaturated fatty acids in loins are of concern because they may cause visual detection of
442 marbling to be more difficult, and loin chops may have a more oily appearance (Johnson, 2008).
443 However, though unsaturated fatty acids would be oilier and less firm than saturated fatty acids,
444 little research has been conducted to understand the relationship between fatty acid composition
445 of pork loin extractable lipid and firmness of pork loins. Of all subjective and objective firmness
446 traits, only initial firmness was significantly but inversely correlated with iodine value (IV; $P =$
447 0.02 ; $r = -0.199$; Table 7) such that as iodine value increased (fatty acids are more unsaturated),
448 firmness score decreased. Initial subjective firmness measures were not significantly correlated
449 with the total percentage of MUFAs or PUFAs ($P \geq 0.13$), however, initial subjective firmness
450 was positively correlated with total percentage of SFAs ($P < 0.01$; $r = 0.231$), negatively
451 correlated with unsaturated:saturated fatty acid ratio ($P < 0.01$; $r = -0.231$) and tended to
452 negatively correlate with the ratio of PUFA:SFA ($P = 0.06$; $r = -0.154$), all in agreement with
453 observations of correlations between initial firmness and IV. The variation in subjective cut
454 surface firmness measures at 28 d postmortem can be explained partially by unsaturated fatty
455 acid content; 4.4% by total percentage of MUFAs ($P = 0.01$) and 3.1% ($P = 0.03$) by total
456 percentage of PUFAs. Further, subjective cut surface firmness measures tended to be correlated
457 with SFA ($P = 0.06$; $r = -0.151$) and the ratio of unsaturated:saturated fatty acids ($P = 0.06$; $r =$
458 0.151). Durometer readings on the dorsal region of the loin were correlated with total percentage
459 of MUFAs ($P = 0.01$; $r = 0.203$), PUFAs ($P < 0.01$; $r = -0.238$) and the ratio of PUFA:SFA ($P =$
460 0.04 ; $r = -0.167$). The weak correlation between IV and initial firmness and the lack of
461 correlation between firmness and 28 d aged pork IV could be due to the fact that this population

462 of loins had a low extractable lipid content (0.47% - 4.18%, Table 1) compared to pork bellies or
463 because fatty acid profile is not related with pork loin firmness.

464 ***Conclusions***

465 When loins not achieving export standards are removed from the population, initial (1 d
466 postmortem) firmness was not correlated to aged (28 d postmortem) pork quality. Given the
467 lack of correlation between firmness and sensory characteristics, selecting only the firmest loins
468 of a population will likely not improve eating quality. Further, at 28 d postmortem, firmness
469 does explain a small portion of the variation in quality and sensory characteristics; however these
470 measures are not consistent throughout the entire loin. Inconsistencies among subjective and
471 objective firmness measures suggest that use of the durometer may not be the most appropriate
472 way to evaluate fresh pork firmness. Differences in firmness were not explained by mechanistic
473 measures. Further work is needed to determine the most appropriate way to evaluate quality
474 while maintaining the whole muscle integrity of loins destined for export to Japan and other
475 quality driven markets if better selection criteria to improve the eating quality of loins is desired.

476 **LITERATURE CITED**

- 477 AOCS. 1998. Official Methods and Recommended Practices of the AOCS. 5th ed. Am. Oil.
478 Chem. Soc., Champaign, IL.
- 479 Arkfeld, E.K., Young, J.M., Johnson, R.C., Fedler, C.A., Prusa, K., Patience, J.F., Dekkers,
480 J.C.M., Gabler, N.K., Lonergan, S.M., and Huff-Lonergan, E. 2015. Composition and
481 quality characteristics of carcass from pigs divergently selected for residual feed intake
482 on high- or low-energy diets. J. Anim. Sci. doi 10.2527/jas2014-8546
- 483 Boler, D.D., Dilger, A.C., Bidner, B.S., Carr, S.N., Eggert, J.M., Day, J.W., Ellis, M., McKeith,
484 F.K, and Killefer, J. 2010. Ultimate pH explains variation in pork quality traits. J.
485 Muscle Foods. 21:119-130.

- 486 Caine, W.R., Aalhus, J.L, Best, D.R., Dugan, M.E.R., and Jeremiah, L.E. 2003. Relationship of
487 texture profile analysis and Warner-Bratzler shear force with sensory characteristics of
488 beef rib steaks. *Meat Sci.* 64:333-339.
- 489 Davis, G.W., Smith, G.C., Carpenter, Z.L., and Cross, H.R. 1975. Relationships of quality
490 indicators to palatability attributes of pork loins. *J. Anim. Sci.* 41:1305-1313.
- 491 Folch, J., Lees, M. and Slone Stanley, G.H. 1957. A simple method for the isolation and
492 purification of total lipids from animal tissues. *The Journal of Biological Chemistry*,
493 226:497-509.
- 494 Hill, F. 1966. The solubility of intramuscular collagen in meat animals of various ages. *J. Food*
495 *Sci.* 31:161-166.
- 496 Huff-Lonergan, E., Baas, T.J., Malek, M., Dekkers, J.C.M., Prusa, K., and Rothschild, M.F.
497 2002. Correlations among selected pork quality traits *J. Anim. Sci.* 80:617-627.
- 498 Huff-Lonergan, E. and Lonergan, S.M. 2005. Mechanisms of water-holding capacity of meat:
499 The role of postmortem biochemical and structural changes. *Meat Sci.* 71:194-204.
- 500 Johnson, R.C. 2008. Impact of biofuel co-products on carcass quality. In P.C. Garnsworthy and J.
501 Wiseman (Eds.), *Recent advances in animal nutrition* (pp. 349-354). Nottingham
502 University Press, Nottingham, UK.
- 503 Masker, C. 2015. 2014 pork exports top \$6 billion. *Pork Checkoff Report*, 34:1.
504 <https://www.pork.org/checkoff-reports/challenge/2014-pork-exports-top-6-billion/>.
505 Accessed April 27, 2015.
- 506 Melody, J.L, Lonergan, S.M., Rowe, L.J., Huiatt, T.W., Mayes, M.S. and Huff-Lonergan, E.
507 2004. Early postmortem biochemical factors influence tenderness and water-holding
508 capacity of three porcine muscles. *J. Anim. Sci.* 82:1195-1205
- 509 Murphy, R.G.L., Howard, S.T., Woerner, D.R., Pendell, D.L., Dixon, C.L., Desimone, T.L.,
510 Green, M.D., Igo, J.L., Tatum, J.D., and Belk, K.E. 2015. Definition, willingness-to-pay,
511 and ranking of quality attributed of U.S. pork as defined by importers in Asia and
512 Mexico. *J. Anim. Sci.* 93:433-441.
- 513 Novakofski, J., Park, S., Bechtel, P. J., and McKeith, F. K. 1989. Composition of cooked pork
514 chops – Effect of removing subcutaneous fat before cooking. *Journal of Food Science*,
515 54:15-17.
- 516 NPPC. 1991. Procedures to evaluate market hogs (3rd ed.).
- 517 NPPC. 1999. Pork Quality Standards.
- 518 Rincker, P.J., Killefer, J., and McKeith, F.K. 2007. Evaluating and objective to measure fresh
519 pork loin firmness. *Meat Sci.* 77:213-219.
- 520 Seman, D.L., Barron, W.N.G., and Matzinger, M. 2013. Evaluating the ability to measure pork
521 fat quality for the production of commercial bacon. *Meat Sci.* 94:262-266.

- 522 Smith, R.M., Gabler, N.K., Young, J.M., Cai, W., Boddicker, N.J., Anderson, M.J., Huff-
523 Lonergan, E., Dekkers, J.C.M., and Lonergan, S.M. 2011. Effects of selection for
524 decreased residual feed intake on composition and quality of fresh pork. *J. Anim. Sci.*
525 89:192-200.
- 526 Swatland, H.J. 1998. Pork softness assessed subjectively and objectively using vacuum-induced
527 changed in reflectance. *J. Muscle Foods.* 9:339-349.
- 528 Trusell, K.A., Apple, J.K., Yancey, J.W.S., Johnson, T.M., Galloway, D.L., and Stackhouse, R.J.
529 2011. Composition and instrumental firmness variations within fresh pork bellies. *Meat*
530 *Sci.* 88:472-480.
- 531 Waylan, A.T., Unruth, J.A., and Johnson, R.C. 1998. Influence of chop location on boneless pork
532 loin quality. In proceedings of the Kansas State University Swine Day. [http://www.asi.k-](http://www.asi.k-state.edu/doc/swine-day-1998/srp819.pdf)
533 [state.edu/doc/swine-day-1998/srp819.pdf](http://www.asi.k-state.edu/doc/swine-day-1998/srp819.pdf). Accessed January 5, 2014.
- 534 Wheeler, T.L., Shackelford, S.D., and Koohmaraie, M. 2000. Variation in proteolysis, sarcomere
535 length, collagen content, and tenderness among major pork muscles. *J. Anim. Sci.*
536 78:958-965.
- 537

Table 1. Characteristics of loins used in the experiment

Variable	N	Mean	Std Dev	Median	Minimum	Maximum
Back Fat, mm	135	17.12	3.86	17	8	28
Loin Depth, mm	135	55.13	7.82	56	36	73
Percent Lean, %	135	52.88	2.37	53.1	46.77	58.57
HCW, kg	152	89.42	6.82	90.29	71.84	107.48
1 d Color	151	3.20	0.30	3	2.5	4
1 d Marbling	151	1.57	0.57	1.5	1	3
1 d Firmness	151	3.19	0.55	3	2	5
1 d L*	151	44.29	2.32	44.45	38.52	52.08
1 d a*	151	6.31	1.01	6.16	4.17	8.79
1 d b*	151	0.14	0.74	0.11	-1.75	2.86
Ultimate pH, 24 h	151	5.57	0.05	5.56	5.45	5.75
28 d Loin pH	154	5.55	0.07	5.55	5.37	5.74
1 d Jowl IV	149	67.63	2.53	67.20	61.00	75.00
1 d Belly IV	151	64.78	3.68	64.90	55.30	74.60
28 d Purge, %	153	2.86	1.45	2.64	0.19	8.11
28 d Drip Loss, %	154	0.78	0.23	0.75	0.37	1.42
Length, cm	154	60.97	2.38	60.96	54.61	66.29
Flop, cm	154	18.46	2.50	18.54	7.11	23.88
Circumference, cm	154	29.37	1.63	29.53	20.73	33.93
NPPC Pre-Bloom Color	154	2.56	0.50	3	2	3
Japan Pre-Bloom Color	154	2.56	0.50	3	2	3
Aged Loin Marbling	154	1.43	0.55	1	1	3
NPPC Post-Bloom Color	154	2.73	0.47	3	2	4
Japan Post-Bloom Color	154	2.73	0.47	3	2	4
10th Rib Subjective Firmness	154	3.05	0.49	3	2	4
Anterior Subjective Firmness	154	3.03	0.51	3	2	4
Posterior Subjective Firmness	154	3.05	0.51	3	2	4
Cut Firmness	154	2.87	0.52	3	2	4
WBSF Chop Subjective Firmness, kg	154	2.63	0.48	2.58	1.61	5.54
Texture Chop Subjective Firmness	154	2.94	0.47	3	2	4
Sensory Chop Subjective Firmness	154	2.99	0.47	3	2	4
Dorsal Durometer	154	53.84	6.30	54.35	36.6	67.9
Ventral Durometer	154	56.03	8.70	55.45	35.6	88.7
Average Whole Loin Durometer	154	54.93	5.98	55.01	40.81	73.10
WBSF Chop Durometer	154	39.45	6.69	39.35	21.6	55.5
Texture Chop Durometer	154	41.18	7.50	41.15	23.5	58
Sensory Chop Durometer	154	42.09	7.12	42.3	20.2	59.9

28 d WBSF, kg	154	2.63	0.48	2.58	1.61	5.54
28 d Moisture, %	154	0.74	0.01	0.74	0.72	0.75
28 d Extractable Lipid, %	154	1.77	0.62	1.69	0.47	4.18
28 d Protein, %	154	21.71	0.43	21.72	20.42	22.74
28 d Sensory Tenderness	154	8.82	1.11	8.93	5.65	11.98
28 d Sensory Chewiness	154	7.26	1.15	7.13	4.33	9.82
28 d Sensory Juiciness	154	7.36	1.00	7.28	5.03	9.75
28 d Sensory Off-Flavor	154	0.01	0.03	0	0	0.24
Soluble Collagen, µg/gram of meat	154	3.95	2.23	3.51	0.65	10.71
Insoluble Collagen, µg/gram of meat	154	23.69	7.75	23.41	2.38	38.24
28 d Cook Loss, %	150	21.37	4.15	21.86	10.96	34.26
Pre-Bloom L*	154	51.22	2.08	51.43	46.23	56.53
Pre-Bloom a*	154	6.64	1.09	6.56	4.1	9.06
Pre-Bloom b*	154	0.19	0.61	0.18	-1.47	1.68
Post-Bloom L*	154	51.36	2.16	51.48	44.47	56.3
Post Bloom a*	154	7.73	1.23	7.7	4.01	10.83
Post Bloom b*	154	3.19	0.81	3.17	0.10	5.14
C 14:0, %	150	1.23	0.12	1.22	0.89	1.61
C 14:1, %	150	0.01	0.02	0	0	0.07
C 15:0, %	150	0.10	0.04	0.09	0	0.23
C 16:0, %	150	21.44	2.64	22.16	10.05	27.52
C 16:1, %	150	3.07	0.57	3.08	1.52	4.54
C 17:0, %	150	0.26	0.07	0.25	0.09	0.48
C 17:1, %	150	0.01	0.08	0	0	0.672
C 18:0, %	150	12.05	2.15	12.30	5.27	16.28
C 18:1 n9, %	150	43.57	5.26	43.64	27.82	56.97
C 18:2 n6, %	150	12.99	2.61	12.83	7.93	21.95
C 18:3 n6, %	150	0.09	0.03	0.09	0	0.17
C 18:3 n3, %	150	0.24	0.04	0.23	0.16	0.41
C 20:0, %	150	0.12	0.03	0.13	0	0.18
C 20:1 n9, %	150	0.51	0.09	0.51	0.31	0.77
C 20:2 n6, %	150	0.28	0.05	0.27	0.17	0.44
C 20:3 n6, %	150	0.37	0.10	0.37	0.14	0.69
C 20:4 n6, %	150	2.83	0.81	2.82	0.50	5.36
C 20:3 n3, %	150	0.05	0.38	0	0	3.965
C 20:5 n3, %	150	0.07	0.03	0.07	0	0.16
C 22:4 n6, %	150	0.41	0.11	0.40	0.18	0.82
C 22:5 n3, %	150	0.24	0.07	0.23	0.08	0.47
C 22:6 n3, %	150	0.08	0.05	0.08	0	0.22
SFA, %	150	35.20	4.63	36.18	16.69	43.58

MUFA, %	150	47.17	5.75	47.31	30.34	62.13
PUFA, %	150	17.63	3.65	17.39	10.15	29.96
UFA:SFA	150	1.91	0.51	1.76	1.29	4.99
PUFA:SFA	150	0.51	0.16	0.49	0.27	1.47
AOCS IV	150	64.14	4.47	63.21	54.51	84.03

538

539

Table 2. Spearman correlation coefficients of firmness traits with other firmness traits

	Subjective						Objective					
	1 d	10th	Anterior	Posterior	Cut Surface	WBSF Chop	Sensory Chop	Dorsal ¹	Ventral ¹	Avg. Durometer ²	WBSF Chop	Sensory Chop
	--	0.141	0.101	0.121	0.195	0.242	0.113	0.126	0.110	0.124	0.057	0.146
1 d		0.08	0.22	0.14	0.02	<0.01	0.17	0.12	0.18	0.13	0.49	0.07
10 th		--	0.620	0.695	0.156	0.096	0.055	0.102	0.068	0.082	-0.132	-0.095
			<0.0001	<0.0001	0.05	0.23	0.50	0.21	0.41	0.31	0.10	0.24
Anterior			--	0.494	0.062	0.151	0.054	0.069	0.043	0.051	-0.042	-0.074
				<0.0001	0.44	0.06	0.51	0.40	0.60	0.53	0.60	0.36
Posterior				--	0.154	0.123	0.053	0.142	0.151	0.170	-0.149	-0.040
					0.06	0.13	0.52	0.08	0.06	0.04	0.06	0.62
Cut Surface					--	0.202	0.147	0.197	0.140	0.201	0.179	0.184
						0.01	0.07	0.01	0.08	0.01	0.03	0.02
WBSF Chop						--	0.484	0.112	0.116	0.144	0.325	0.050
							<0.0001	0.17	0.15	0.07	<0.0001	0.54
Sensory Chop							--	0.082	0.042	0.081	0.141	0.085
								0.31	0.60	0.32	0.08	0.29
Dorsal								--	0.266	0.721	-0.036	0.034
									<0.01	<0.0001	0.66	0.68
Ventral									--	0.844	0.036	0.081
										<0.0001	0.66	0.32
Avg. Durometer ²										--	0.005	0.078
											0.95	0.34
WBSF Chop											--	0.401
												<0.0001

540 ¹ Durometer readings were determined at approximately the 10th rib on both the dorsal and ventral sides of the loin.

541 ² An average whole loin durometer reading was calculated as the average of the durometer readings on the dorsal and ventral sides of
542 the loin.

Table 3. Spearman correlation coefficients of firmness traits with 28 d aged meat quality

	Subjective							Objective				
	1 d	10th	Anterior	Posterior	Cut Surface	WBSF Chop	Sensory Chop	Dorsal ¹	Ventral ¹	Avg. Durometer ²	WBSF Chop	Sensory Chop
Loin Length	-0.022	-0.028	-0.232	-0.043	0.026	-0.050	-0.028	-0.053	0.104	0.039	0.091	0.049
	0.79	0.73	<0.01	0.59	0.75	0.53	0.73	0.51	0.20	0.63	0.26	0.54
	0.443	0.268	0.216	0.222	0.112	0.049	0.019	0.076	0.034	0.049	-0.027	0.064
Loin Flop	<0.0001	<0.01	0.01	0.01	0.17	0.54	0.82	0.35	0.68	0.55	0.74	0.43
NPPC Color Pre-Bloom	0.065	0.001	0.071	-0.068	0.147	0.227	0.115	0.032	0.080	0.102	0.148	0.192
	0.43	0.99	0.38	0.40	0.07	<0.01	0.16	0.69	0.32	0.21	0.07	0.02
Japan Color Pre-Bloom	0.065	0.001	0.071	-0.068	0.147	0.227	0.115	0.032	0.080	0.102	0.148	0.192
	0.43	0.99	0.38	0.40	0.07	<0.01	0.16	0.69	0.32	0.21	0.07	0.02
NPPC Color Post-Bloom	0.113	-0.006	0.000	0.002	0.115	0.049	0.014	0.070	0.055	0.106	0.053	0.257
	0.17	0.94	1.00	0.98	0.16	0.54	0.87	0.39	0.50	0.19	0.51	<0.01
Japan Color Post-Bloom	0.113	-0.006	0.000	0.002	0.115	0.049	0.014	0.070	0.055	0.106	0.053	0.257
	0.17	0.94	1.00	0.98	0.16	0.54	0.87	0.39	0.50	0.19	0.51	<0.01
Aged Marbling	0.013	-0.035	-0.030	-0.071	0.166	0.024	-0.005	0.236	0.005	0.125	0.043	-0.075
	0.87	0.66	0.71	0.38	0.04	0.77	0.95	<0.01	0.95	0.12	0.59	0.36
Moisture	-0.025	0.036	0.062	0.028	-0.096	0.030	-0.090	-0.109	0.093	-0.0001	-0.101	-0.091
	0.76	0.66	0.45	0.73	0.23	0.72	0.26	0.18	0.25	1.00	0.21	0.26
Extractable Lipid	0.071	-0.080	-0.128	-0.059	0.138	-0.042	0.151	0.165	-0.071	0.034	0.178	0.210
	0.39	0.32	0.11	0.47	0.09	0.61	0.06	0.04	0.38	0.67	0.03	0.01
Protein, %	0.013	0.099	0.070	0.048	0.004	0.010	-0.122	-0.056	-0.045	-0.054	-0.154	-0.046
	0.87	0.22	0.39	0.56	0.96	0.90	0.13	0.49	0.58	0.51	0.06	0.57
L* Pre-Bloom	-0.137	-0.056	-0.095	-0.099	-0.088	-0.023	-0.039	-0.166	-0.030	-0.125	0.094	-0.117
	0.09	0.49	0.24	0.22	0.28	0.77	0.63	0.04	0.71	0.12	0.25	0.15
a* Pre-Bloom	0.050	0.145	0.054	0.157	0.175	0.067	0.044	0.042	0.087	0.102	-0.023	0.099
	0.54	0.07	0.50	0.05	0.03	0.41	0.59	0.61	0.28	0.21	0.78	0.22

	-0.122	0.092	0.013	0.043	0.086	-0.071	-0.075	-0.136	0.056	-0.024	0.062	-0.074
b* Pre-Bloom	0.14	0.26	0.87	0.59	0.29	0.38	0.35	0.09	0.49	0.77	0.45	0.36
	-0.101	-0.061	-0.042	-0.067	-0.059	-0.074	-0.009	-0.111	0.044	-0.035	0.000	-0.069
L* Post-Bloom	0.22	0.45	0.60	0.41	0.47	0.36	0.92	0.17	0.59	0.66	1.00	0.39
	0.036	0.123	0.084	0.069	0.181	0.039	0.088	0.020	0.080	0.074	0.058	0.090
a* Post-Bloom	0.66	0.13	0.30	0.39	0.02	0.63	0.28	0.80	0.33	0.36	0.47	0.27
	-0.068	0.104	0.033	0.063	0.116	-0.064	0.132	-0.027	0.138	0.088	0.015	0.014
b* Post-Bloom	0.41	0.20	0.69	0.44	0.15	0.43	0.10	0.74	0.09	0.28	0.85	0.87

543 ¹ Durometer were determined at approximately the 10th rib on both the dorsal and ventral sides of the loin.

544 ² An average whole loin durometer reading was calculated as the average of the durometer readings on the dorsal and ventral sides of
545 the loin.

546

Table 4. Spearman correlation coefficients of firmness with pH, WHC measures, sensory characteristics, and collagen content

	Subjective							Objective				
	1 d	10th	Anterior	Posterior	Cut Surface	WBSF Chop	Sensory Chop	Dorsal ¹	Ventral ¹	Avg. Durometer ²	WBSF Chop	Sensory Chop
	0.174	-0.074	-0.005	-0.053	0.071	0.211	0.016	0.031	0.142	0.135	0.131	0.164
WBSF	0.03	0.36	0.95	0.52	0.38	0.01	0.85	0.70	0.08	0.09	0.11	0.04
Tenderness	-0.022	0.129	0.170	0.008	-0.140	-0.076	-0.164	0.009	0.018	-0.011	-0.022	-0.134
	0.79	0.11	0.03	0.92	0.08	0.35	0.04	0.92	0.83	0.90	0.78	0.10
Chewiness	0.057	-0.180	-0.189	-0.070	-0.028	0.117	0.144	0.072	-0.006	0.056	-0.030	0.119
	0.49	0.03	0.02	0.39	0.73	0.15	0.07	0.37	0.94	0.49	0.71	0.14
Juiciness	0.143	0.070	0.132	0.056	-0.092	0.063	-0.009	-0.190	-0.087	-0.176	0.018	-0.098
	0.08	0.39	0.10	0.49	0.26	0.44	0.91	0.02	0.28	0.03	0.82	0.23
Off Flavor	0.081	-0.021	-0.012	-0.023	-0.063	-0.030	0.007	-0.129	-0.254	-0.229	0.025	0.066
	0.32	0.80	0.88	0.78	0.44	0.71	0.93	0.11	<0.01	<0.01	0.76	0.42
Aged pH	0.058	0.111	0.086	0.111	0.038	0.015	-0.108	0.197	0.104	0.183	-0.123	-0.061
	0.48	0.17	0.29	0.17	0.64	0.86	0.18	0.01	0.20	0.02	0.13	0.46
Purge	0.136	-0.268	-0.349	-0.304	-0.0028	0.101	0.094	0.029	-0.131	-0.068	0.168	0.093
	0.10	<0.01	<0.0001	0.0001	0.73	0.21	0.25	0.72	0.11	0.41	0.04	0.25
Drip Loss	-0.028	-0.100	0.134	-0.039	-0.213	0.027	-0.079	-0.068	-0.083	-0.099	0.036	0.026
	0.73	0.22	0.10	0.63	0.01	0.74	0.33	0.40	0.31	0.22	0.66	0.75
Cook Loss	-0.028	-0.001	0.054	-0.032	0.002	0.103	0.077	-0.076	0.023	-0.018	0.019	0.054
	0.73	0.99	0.51	0.69	0.98	0.21	0.35	0.36	0.78	0.83	0.81	0.51
Soluble Collagen	-0.078	-0.105	-0.047	-0.026	-0.058	0.069	0.144	-0.019	0.011	0.005	0.024	0.052
	0.34	0.19	0.56	0.74	0.47	0.39	0.08	0.81	0.89	0.96	0.76	0.53
Insoluble Collagen	0.060	0.004	0.081	0.036	-0.078	0.220	0.043	-0.117	-0.106	-0.130	0.025	-0.130
	0.47	0.96	0.32	0.66	0.33	0.01	0.60	0.15	0.19	0.11	0.76	0.11

547 ¹ Durometer were determined at approximately the 10th rib on both the dorsal and ventral sides of the loin.

548 ² An average whole loin durometer reading was calculated as the average of the durometer readings on the dorsal and ventral sides of
549 the loin.

Table 5. Spearman correlation coefficients of firmness traits with production facility data

	Subjective							Objective				
	1 d	10th	Anterior	Posterior	Cut Surface	WBSF Chop	Sensory Chop	Dorsal ¹	Ventral ¹	Avg. Durometer ²	WBSF Chop	Sensory Chop
Back Fat, mm	0.158	0.206	0.055	0.167	0.258	-0.167	-0.071	0.145	0.104	0.117	-0.062	0.000
	0.07	0.02	0.53	0.05	<0.01	0.05	0.41	0.09	0.23	0.18	0.48	1.00
Loin Depth, mm	0.182	-0.029	0.088	-0.009	-0.100	-0.017	-0.048	0.156	-0.019	0.077	-0.155	-0.004
	0.04	0.74	0.31	0.92	0.25	0.85	0.58	0.07	0.82	0.36	0.07	0.96
Percent Lean, %	-0.046	-0.152	0.013	-0.143	-0.228	0.041	-0.005	0.047	-0.077	-0.004	-0.081	-0.030
	0.60	0.08	0.89	0.10	0.01	0.63	0.95	0.59	0.37	0.96	0.35	0.73
HCW, kg	0.325	0.150	0.012	0.092	0.127	0.055	0.047	0.061	0.106	0.096	0.057	0.029
	<0.0001	0.06	0.88	0.26	0.12	0.50	0.57	0.45	0.19	0.24	0.49	0.73
Color	0.095	0.080	0.108	0.086	0.149	0.104	0.143	0.175	0.000	0.114	-0.041	0.041
	0.24	0.33	0.19	0.29	0.07	0.20	0.08	0.03	1.00	0.16	0.62	0.62
Marbling	0.060	0.042	0.121	0.077	0.182	0.058	-0.013	0.173	0.140	0.194	0.069	0.005
	0.47	0.61	0.14	0.35	0.03	0.48	0.87	0.03	0.09	0.02	0.40	0.95
L*	-0.006	-0.059	-0.093	-0.020	-0.062	-0.114	-0.118	0.061	0.087	0.096	-0.110	-0.032
	0.94	0.47	0.25	0.81	0.45	0.16	0.15	0.45	0.29	0.24	0.18	0.69
a*	0.058	0.207	0.141	0.139	0.118	0.148	0.127	0.037	-0.072	-0.023	0.072	0.017
	0.48	0.01	0.09	0.09	0.15	0.07	0.12	0.65	0.38	0.76	0.38	0.84
b*	-0.018	0.105	-0.046	0.076	0.060	-0.049	-0.087	0.060	0.123	0.125	0.026	-0.077
	0.83	0.20	0.57	0.35	0.47	0.55	0.29	0.46	0.13	0.13	0.75	0.35
24 h pH	0.040	0.050	0.060	0.082	0.143	0.069	0.042	0.060	-0.003	0.036	-0.031	-0.093
	0.63	0.54	0.46	0.32	0.08	0.40	0.61	0.47	0.97	0.66	0.71	0.26
Jowl IV	-0.127	-0.048	-0.023	-0.018	0.028	0.039	-0.013	0.111	-0.118	-0.017	-0.013	-0.037
	0.12	0.56	0.78	0.83	0.73	0.63	0.88	0.18	0.15	0.83	0.87	0.65
Belly IV	-0.082	-0.053	-0.049	-0.060	0.131	0.037	-0.056	0.073	-0.118	-0.030	0.008	-0.038
	0.32	0.52	0.55	0.46	0.11	0.65	0.49	0.37	0.15	0.72	0.93	0.65

550 ¹ Durometer were determined at approximately the 10th rib on both the dorsal and ventral sides of the loin.

551 ² An average whole loin durometer reading was calculated as the average of the durometer readings on the dorsal and ventral sides of
552 the loin.

553

Table 6. Spearman correlations coefficients of color, marbling, and sensory characteristics

	1 d color score	28 d color score	1 d marbling score	28 d marbling score
	-0.019	-0.059	-0.055	-0.049
Tenderness	0.82	0.47	0.50	0.54
	0.088	0.109	-0.065	-0.041
Chewiness	0.28	0.18	0.43	0.61
	-0.039	0.014	-0.064	-0.028
Juiciness	0.63	0.86	0.43	0.73
	-0.148	-0.010	-0.080	-0.054
Off Flavor	0.07	0.90	0.33	0.51

554

555

Table 7. Spearman correlation coefficients of firmness traits with fatty acid methyl ester profile (FAME) and iodine value (IV)

	Subjective							Objective				
	1 d	10th	Anterior	Posterior	Cut Surface	WBSF Chop	Sensory Chop	Dorsal ¹	Ventral ¹	Avg. Durometer ²	WBSF Chop	Sensory Chop
	-0.015	-0.063	-0.001	-0.004	-0.012	-0.116	-0.004	0.108	0.022	0.096	-0.054	0.034
C 14:0, %	0.86	0.44	0.99	0.96	0.88	0.16	0.96	0.19	0.79	0.24	0.52	0.68
	-0.140	0.079	0.033	0.154	0.052	0.041	0.099	0.017	0.035	0.034	-0.053	-0.092
C 14:1, %	0.09	0.34	0.69	0.06	0.53	0.62	0.23	0.84	0.67	0.68	0.52	0.26
	0.035	-0.118	-0.073	-0.078	-0.051	0.099	-0.005	-0.073	0.065	-0.007	0.052	0.045
C 15:0, %	0.67	0.15	0.38	0.34	0.54	0.23	0.95	0.38	0.43	0.93	0.53	0.58
	0.214	-0.004	-0.057	0.091	-0.103	0.017	-0.025	-0.014	0.049	0.017	-0.049	-0.031
C 16:0, %	0.01	0.96	0.49	0.27	0.21	0.84	0.76	0.87	0.55	0.83	0.55	0.71
	-0.127	0.093	0.062	0.042	0.189	-0.065	-0.003	0.252	0.022	0.163	0.001	0.010
C 16:1, %	0.13	0.26	0.45	0.61	0.02	0.43	0.97	<0.01	0.79	0.05	0.99	0.90
	0.030	-0.162	-0.164	-0.091	-0.198	-0.003	-0.044	-0.148	0.035	-0.085	-0.011	-0.135
C 17:0, %	0.72	0.05	0.04	0.27	0.01	0.97	0.59	0.07	0.67	0.30	0.90	0.10
	0.134	-0.018	-0.010	0.062	-0.120	0.078	0.011	-0.078	-0.130	-0.132	-0.055	-0.048
C 17:1, %	0.11	0.83	0.90	0.45	0.15	0.34	0.90	0.34	0.11	0.11	0.51	0.56
	0.204	-0.095	-0.073	-0.031	-0.173	0.020	-0.034	-0.160	0.056	-0.064	-0.029	-0.030
C 18:0, %	0.01	0.25	0.38	0.71	0.03	0.81	0.68	0.05	0.49	0.44	0.73	0.71
C 18:1 n9, %	-0.122	0.069	0.073	-0.033	0.202	0.006	0.095	0.197	-0.085	0.059	0.056	0.081
	0.14	0.40	0.38	0.69	0.01	0.94	0.25	0.02	0.30	0.47	0.49	0.33
C 18:2 n6, %	-0.108	-0.028	-0.017	-0.009	-0.221	-0.051	-0.090	-0.256	-0.042	-0.163	-0.036	0.016
	0.19	0.74	0.84	0.92	<0.01	0.53	0.27	<0.01	0.61	0.05	0.66	0.85
C 18:3 n6, %	-0.045	-0.043	-0.093	-0.052	0.078	0.008	-0.073	-0.068	0.006	-0.051	0.024	0.014
	0.59	0.60	0.26	0.53	0.34	0.92	0.37	0.41	0.95	0.54	0.77	0.86
C 18:3 n3, %	-0.150	-0.028	0.059	-0.046	-0.181	0.016	-0.038	-0.190	-0.138	-0.194	0.011	0.008
	0.07	0.74	0.47	0.58	0.03	0.85	0.64	0.02	0.09	0.02	0.89	0.92

	0.199	-0.076	0.002	-0.019	-0.165	0.065	-0.011	0.007	0.019	0.004	0.012	-0.020
C 20:0, %	0.02	0.36	0.98	0.81	0.04	0.43	0.89	0.93	0.81	0.95	0.89	0.81
C 20:1 n9, %	-0.032	-0.042	-0.008	-0.075	0.121	0.074	0.108	0.083	-0.041	0.024	0.087	0.072
	0.70	0.61	0.92	0.36	0.14	0.37	0.19	0.31	0.62	0.77	0.29	0.38
C 20:2 n6, %	-0.200	-0.113	-0.059	-0.122	-0.209	0.060	0.049	-0.222	-0.125	-0.193	0.043	0.047
	0.02	0.17	0.48	0.14	0.01	0.47	0.55	<0.01	0.13	0.02	0.60	0.57
C 20:3 n6, %	-0.077	0.008	-0.055	-0.011	-0.111	0.005	-0.030	-0.168	-0.036	-0.120	-0.031	-0.014
	0.36	0.92	0.50	0.89	0.18	0.95	0.72	0.04	0.66	0.14	0.71	0.87
C 20:4 n6, %	-0.102	0.003	-0.099	-0.009	-0.061	0.005	-0.040	-0.191	-0.046	-0.141	-0.020	-0.052
	0.22	0.97	0.23	0.92	0.45	0.95	0.62	0.02	0.58	0.09	0.81	0.53
C 20:3 n3, %	-0.115	-0.081	-0.068	-0.058	0.007	0.073	0.089	0.050	-0.107	-0.024	-0.054	0.036
	0.17	0.32	0.41	0.48	0.93	0.37	0.28	0.54	0.19	0.77	0.51	0.66
C 20:5 n3, %	-0.012	-0.051	-0.086	-0.051	-0.040	0.061	-0.037	-0.107	0.045	-0.036	0.026	0.009
	0.89	0.54	0.30	0.53	0.63	0.46	0.66	0.19	0.58	0.67	0.76	0.92
C 22:4 n6, %	-0.069	-0.012	-0.138	0.002	-0.054	0.029	-0.056	-0.175	-0.040	-0.117	-0.006	-0.037
	0.41	0.89	0.09	0.98	0.51	0.73	0.49	0.03	0.62	0.15	0.95	0.65
C 22:5 n3, %	-0.043	-0.032	-0.098	-0.019	-0.096	0.005	-0.061	-0.200	-0.036	-0.127	-0.041	-0.011
	0.60	0.70	0.23	0.82	0.24	0.95	0.45	0.01	0.66	0.12	0.62	0.89
C 22:6 n3, %	-0.051	-0.043	-0.058	-0.027	-0.026	0.168	0.058	-0.056	-0.045	-0.037	0.023	-0.004
	0.54	0.60	0.48	0.74	0.75	0.04	0.48	0.49	0.58	0.66	0.78	0.96
SFA, %	0.231	-0.062	-0.074	0.028	-0.151	0.028	-0.026	-0.092	0.049	-0.029	-0.024	0.038
	<0.01	0.45	0.37	0.73	0.06	0.74	0.75	0.26	0.55	0.72	0.77	0.65
MUFA, %	-0.125	0.070	0.072	-0.027	0.209	0.005	0.096	0.203	-0.079	0.066	0.058	0.054
	0.13	0.39	0.38	0.74	0.01	0.96	0.24	0.01	0.34	0.42	0.48	0.51
PUFA, %	-0.107	-0.024	-0.038	-0.011	-0.175	-0.018	-0.062	-0.238	-0.044	-0.153	-0.160	-0.042
	0.20	0.77	0.65	0.90	0.03	0.83	0.45	<0.01	0.59	0.06	0.05	0.61
Ratio Unsaturated FA:SFA	-0.231	0.062	0.074	-0.028	0.151	-0.028	0.026	0.092	-0.049	0.029	0.024	-0.038
	<0.01	0.45	0.37	0.73	0.06	0.74	0.75	0.26	0.55	0.72	0.77	0.65
Ratio	-0.154	0.006	-0.008	-0.008	-0.083	-0.029	-0.051	-0.167	-0.073	-0.132	-0.151	-0.036

PUFA:SFA	0.06	0.94	0.92	0.92	0.31	0.73	0.53	0.04	0.38	0.11	0.06	0.66
	-0.199	0.045	0.079	-0.015	0.016	-0.043	-0.010	-0.023	-0.065	-0.047	0.021	0.055
AOCS IV	0.02	0.59	0.33	0.85	0.84	0.60	0.91	0.78	0.43	0.57	0.79	0.51

556 ¹ Durometer were determined at approximately the 10th rib on both the dorsal and ventral sides of the loin.

557 ² An average whole loin durometer reading was calculated as the average of the durometer readings on the dorsal and ventral sides of
 558 the loin.

559

560