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Effect of the Programmed Nutrition Beef Program on moisture retention of cooked ground beef patties and enhanced strip loins^{1,2}

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15 **ABSTRACT:**

16 This study evaluated the influence of the Programmed Nutrition Beef Program and exogenous
17 growth promotants (**ExGP**) on water holding capacity characteristics of enhanced beef strip
18 loins. Sixty, frozen strip loins, arranged in a 2×2 factorial treatment arrangement with dietary
19 program serving as the first factor and use of ExGP as the second factor, were thawed, injected
20 with an enhancement solution, and stored for 7 d. Loins from ExGP cattle possessed the ability
21 to bind more ($P < 0.05$) water before pumping and bind less ($P < 0.05$) water after pumping and
22 storage. Loin pH across treatments was similar ($P > 0.10$) before injection, but increased post-
23 injection and after storage ($P < 0.01$). Treatments did not affect loin purge loss, steak cook loss,
24 and expressible moisture ($P > 0.10$). The Programmed Nutrition Beef Program and use of ExGPs
25 minimally impacted water holding capacity of enhanced frozen/thawed beef strip loins.

26 **Keywords:** Water-holding; Beef; Ractopamine; Feeding, Enhancement

27 **1. Introduction**

28 The goal of the beef industry is to produce a consistent, high quality product as efficiently
29 as possible. Many different feedlot management strategies are employed to maximize efficiency
30 including the utilization of different feed additives such as monensin (Rumensin; Elanco Animal
31 Health, Greenfield IN) and tylosin (Tylan; Elanco Animal Health), and growth promoting
32 technologies such as beta-adrenergic agonists and steroidal implants. Utilization of feed
33 additives, implants, and beta-adrenergic agonists can greatly improve efficiency of beef
34 production, but also can have negative consequences for beef quality (Winterholler et al., 2008).
35 Beef palatability encompasses characteristics such as juiciness, tenderness, and flavor (Platter,
36 Tatum, Belk, Scanga, & Smith, 2003), but water-holding capacity also impacts beef palatability.
37 The ability of whole muscle or ground beef to retain moisture during storage and through
38 processing is an important quality characteristic. It is established moisture loss from whole
39 muscle products can impact the amount of salable product (Offer et al., 1989; Huff-Lonergan and
40 Lonergan, 2005). Although cook yields of ground beef can be mostly dependent on fat content
41 of the blend (Roth, McKeith, & Brewer, 1999), the ability of product to retain water during
42 processing and cooking could also be important.

43 The Programmed Nutrition Beef Program (Alltech Inc., Nicholasville, KY) consists of
44 two products that are designed to replace components of conventional feedlot diets. The
45 Programmed Nutrition Beef Receiver is intended to be fed during the step-up period at the rate of
46 $14 \text{ g}\cdot\text{animal}^{-1}\cdot\text{d}^{-1}$, while Programmed Nutrition Beef Finisher is fed for the remainder of the
47 finishing period at a rate of $20 \text{ g}\cdot\text{animal}^{-1}\cdot\text{d}^{-1}$. Previous research by Phelps et al. (2014) reported
48 the use of Programmed Nutrition Beef Program supplements in finishing feedlot diets decreased
49 purge loss of strip loins aged 14 d compared to strip loins from steers fed a diet containing

50 conventional feedlot dietary components. Additionally, steaks from steers fed the Programmed
51 Nutrition Beef Program supplements and steaks from steers that were not administered
52 exogenous growth promotants (**ExGP**) had a decreased cook loss when compared to steaks from
53 the conventional feedlot diet and administered ExGP, respectively. The objectives of this study
54 were to evaluate influence of the Programmed Nutrition Beef Program and ExGPs on moisture
55 retention of 85% lean ground beef patties, brine uptake and retention, objective water holding
56 capacity, and Warner-Bratzler shear force (**WBSF**) of strip loins that were thawed from a frozen
57 state and enhanced 110% with a salt/tripolyphosphate solution.

58 **2. Materials and Methods**

59 *2.1 Animals*

60 Crossbred feedlot steers (initial BW 383 kg \pm 30; n = 64 pens; 16 pens/treatment; 8
61 steers/pen) were blocked by body weight and subjected to a randomized complete block design
62 with 2 \times 2 factorial treatment arrangement. Steers were separated into a conventional finishing
63 program treatment (**CON**) or Alltech Programmed Nutrition Beef Program treatment (**PN**; Table
64 1). Conventional diets contained monensin, tylosin, vitamins A and E, and supplemented trace
65 minerals including copper sulfate, cobalt carbonate, ethlenediamine dihydriodide, manganous
66 sulfate sodium selenite, and zinc sulfate. The Programmed Nutrition Beef Receiver portion of the
67 diet was included in the total mixed ration for the first 21 d at a rate of 14 g \cdot animal⁻¹ \cdot d⁻¹. The
68 Programmed Nutrition Beef Finisher was included in the total mixed ration at a rate of 20
69 g \cdot animal⁻¹ \cdot d⁻¹ for the final 154 d of the feeding period. The components of the Alltech
70 Programmed Nutrition Beef Program diet were premixed into a ground corn carrier and
71 subsequently blended into the total mixed ration. Both the Programmed Nutrition Beef Receiver
72 and Finisher supplements contained a proprietary blend of organic trace elements, ascorbic acid,

73 *Lactobacillus acidophilus* fermentation product, *Enterococcus faecium* fermentation product, and
74 selenium yeast. Additionally, Programmed Nutrition Beef Receiver included *Aspergillus niger*
75 fermentation extract and Programmed Nutrition Beef Finisher included *Aspergillus oryzae*
76 fermentation extract. Each diet was fed in conjunction with (**EGP+**) or in the absence of (**EGP-**)
77 exogenous growth promotants (**ExGP**). Steers receiving **ExGP** were administered a Component
78 E-S implant on d 1 of the study, reimplanted with Component TE-IS (Elanco Animal Health) on
79 d 94, and fed ractopamine hydrochloride (**RAC**; Elanco Animal Health) at a rate of $400 \cdot \text{d}^{-1} \cdot \text{steer}^{-1}$
80 the final 28 d before harvest.

81 2.2 Loin collection

82 On d 175 of the study, animals were shipped 430 km to a commercial abattoir (Tyson
83 Fresh Meats, Holcomb, KS) for harvest. After a 36-h chill, strip loins (Institutional Meat
84 Purchase Specifications 180) were removed from the left side of 2 carcasses selected at random
85 from each pen and were transported back to the Kansas State University Meats Laboratory.
86 Loins were wet-aged for 14 d, 5 steaks were fabricated from the anterior portion of the
87 *Longissimus lumborum* for another study (Phelps et al., 2014), and the remainder of the
88 *Longissimus lumborum* was frozen at $-40\text{ }^{\circ}\text{C}$.

89 2.3 Loin Selection and Sampling

90 The posterior portion of the *Longissimus lumborum* of sixty frozen beef strip loins (n =
91 15 per treatment) were selected at random from the previous study (Phelps et al., 2014). For the
92 injection study, loins selected were large enough to be injected (smallest loin = 0.58 kg), and also
93 yielded 500 g of tissue for grinding. This resulted in a sample distribution that represented all 16
94 of the original weight blocks and 52 of the pens of the Phelps et al. (2014) study.

95 2.4 Loin processing and injection

96 Forty-eight hours prior to processing, strip loins were removed from the freezer and
97 thawed at 2 ± 1 °C. Once thawed, all subcutaneous fat was removed from each loin, identified
98 by loin number, and reserved for subsequent preparation of ground beef patties. From each
99 trimmed loin, approximately 500 g of muscle was removed for grinding from the most anterior
100 portion of each loin. Additionally, a 120-g sample was removed for measurement of expressible
101 moisture (**EM**) and water binding ability (**WBA**) as described below. Before injection, pH was
102 measured using a pH probe designed for use in meat (Model HI 99163; Hanna Instruments,
103 Smithfield, RI) and loin sections were weighed. Loin sections were injected using a multi-needle
104 injector (Model N50: Schröder Maschinenbau GmbH, Werther, Germany) with a brine solution
105 (pH 7.95) which contained 5% sodium chloride and 3% sodium tripolyphosphate (Brifisol 85
106 Instant; BK Giulini Corp., Simi Valley, CA) to achieve a 0.5% sodium chloride and 0.3%
107 sodium tripolyphosphate level within the muscle when pumped 10% over green weight.
108 Following a 2-min rest period, loins were reweighed to ensure brine was assimilated to achieve
109 110% above green weight (average $111.14 \pm 1.46\%$) and pH was recorded again before injected
110 loins were vacuum packaged in 3-mm high-barrier vacuum pouches (Prime Source, Bunzyl
111 Processor Division, Kansas City, MO). After storage for 7 d at 2 ± 1 °C, loins were removed
112 from packages, patted dry, and re-weighed for purge loss calculations [(injected weight-stored
113 weight)/injected weight] $\times 100$. At this time, pH was measured, a 120-g sample was removed for
114 expressible moisture and WBA analyses, and a 2.54-cm thick steak was fabricated for
115 determination of WBSF.

116 *2.5 Grinding*

117 To formulate a ground beef blend that was 85% lean and 15% fat, analyzed fat
118 percentages from proximate analysis were used to calculate a blend for each sample. Using a

119 table top grinder (Model KG-12-FS; Pro-Cut, Houston, TX), lean and fat for each sample were
120 ground separately through a 9.5-mm plate, blended together by hand, and then reground through
121 a 3.2-mm plate. Once ground, two 1.2-cm thick × 10.7-cm wide patties weighing 112 g were
122 formed from each sample.

123 *2.6 Cooking and Warner-Bratzler shear force*

124 All cooking and WBSF procedures were conducted according to the Meat Cookery and
125 Sensory Guidelines (AMSA, 1995). Ground beef patties were weighed before cooking and
126 cooked on a flat top grill (Model 106733; Walmart, Bentonville, AR) set to 132 °C. Patties were
127 turned at 1 min, turned again at 2 min, and then turned every 2 min until they reached an internal
128 temperature of 71 °C. A hypodermic needle thermometer consisting of a copper-constantan probe
129 connected to a Doric Trendicator 410A monitor (VAS Engineering; San Francisco, CA) was
130 used to monitor internal temperature. Patties were cooled at room temperature and once they
131 could be handled they were reweighed to determine cook loss [(precooked weight-cooked
132 weight)/precooked weight] ×100.

133 Prior to cooking, steaks were weighed and a thermocouple wire (30-gauge copper-
134 constantan; Omega Engineering, Stamford, CT) was inserted into the geometric center of each
135 steak to monitor internal temperature using a Doric Minitrend 205 monitor (VAS Engineering).
136 Steaks were cooked on electric, open-hearth Hamilton Beach grills (Indoor/Outdoor; Southern
137 Pines, NC) preheated to 204 °C. Steaks were turned when they reached an internal temperature
138 of 40 °C, removed from grills at 70 °C, and allowed to cool until they could be handled before
139 being reweighed to determine cook loss [(precooked weight-cooked weight)/precooked weight]
140 ×100. Steaks were chilled for 24 h at 7 ± 1 °C, and six 1.27-cm cores were then removed from

141 each steak parallel to the muscle fiber. Cores were sheared once through the center using an
142 INSTRON Model 5569 testing machine (Instron, Canton, MA) with a Warner-Bratzler shear
143 head attached (100-kg compression load cell, crosshead speed of 250 mm/min).

144 *2.7 Water binding ability and expressible moisture*

145 The WBA of samples was measured by protein swelling analysis as described by
146 Pietrasik and Janz (2009) with a modification. The modification made was the 100-g sample of
147 muscle was cut into $0.5 \times 0.5 \times 0.5$ cm pieces and then blended in a Waring blender (Waring
148 Products Division, Hartford, CT) with 300 mL of distilled water. The remainder of the procedure
149 was carried out as described by the authors.

150 Expressible moisture of samples was measured using the centrifugation method described
151 by Jauregui, Regenstein, and Baker (1981). A 5-g sample of muscle was weighed, placed in a 50
152 mL conical tube on top of 25 g of 4-mm glass beads, and centrifuged at $900 \times g$ for 10 min. After
153 centrifugation, samples were removed from tubes and reweighed. Percent EM was calculated as
154 $[(\text{initial weight} - \text{centrifuged weight}) / \text{initial weight}] \times 100$.

155 *2.8 Statistical analyses*

156 Purge loss, cook loss, and WBSF data were analyzed as a randomized complete block
157 design with a 2×2 factorial arrangement. Dietary program and ExGP served as the main effects
158 and the animal weight block from Phelps et al. (2014) was the random effect. For loin pH,
159 expressible moisture, and WBA, data were analyzed as repeated measures using a randomized
160 complete block design with a 2×2 factorial treatment arrangement. Period served as the
161 repeated measure with loin (observational unit) as the subject and compound symmetry as
162 covariance structure. The PROC MIXED procedure of SAS (SAS Inst. Inc., Cary, NC) was

163 utilized and pair-wise comparisons between the least-squares means of the factor levels were
164 computed using the PDIFF option of the LSMEANS statement. Differences were considered
165 significant at $\alpha \leq 0.05$ and tendencies at $\alpha \leq 0.10$.

166 **3. Results**

167 *3.1 Loin pH*

168 Loin pH was recorded at three periods: before injection, after injection, and after 7-d of
169 storage post injection (Table 2). There was no three-way interaction ($P = 0.55$) of dietary
170 program \times ExGP \times period for pH. Also, the two-way interaction of dietary program \times ExGP did
171 not affect ($P = 0.39$) loin pH, but there was a tendency ($P = 0.10$) for a dietary program \times period
172 interaction to affect loin pH. Pre-injection, the pH of CON and PN loins was similar ($P = 0.75$),
173 but post-injection PN loins had a greater ($P = 0.002$) pH than CON loins. Additionally, the post-
174 injection pH of loins from both CON and PN loins increased ($P < 0.01$) by 0.37 and 0.32 units
175 compared to their pre-injection pH, respectively. The pH recorded for both dietary treatments
176 after 7 d of storage was similar to that recorded post-injection ($P > 0.15$); however, there was no
177 difference ($P = 0.50$) in pH between the two dietary treatments after 7 d of storage. Also, pH data
178 indicate an ExGP \times period interaction ($P = 0.02$). Pre-injection pH was similar ($P = 0.88$)
179 between EGP+ and EGP- loins. Post-injection pH of EGP- loins was greater ($P = 0.001$) than
180 EGP+ loins, but pH recorded after 7-d of storage was similar ($P = 0.89$) between the two
181 treatments. The pH of loins from the two dietary treatments differed with PN loins having a pH
182 that was 0.04 units greater ($P = 0.02$) than CON loins. Additionally, there was a tendency ($P =$
183 0.052) for pH difference between the two ExGP treatments. Finally, as expected, period elicited
184 an effect ($P < 0.01$) on the pH of loins. Before injection, the pH of loins was 5.53, and post-

185 injection pH was 0.32 units greater ($P < 0.01$). The pH recorded after 7 d of storage was
186 comparable ($P = 0.77$) to post-injection pH.

187 *3.2 Purge loss, cook loss, and WBSF*

188 For purge loss, steak cook loss, and ground beef cook loss there was no two-way
189 interaction of dietary program \times ExGP (Table 3; $P > 0.34$). Also, the main effect of dietary
190 program did not affect purge loss, steak cook loss, and ground beef cook loss ($P > 0.35$).
191 Additionally, the ExGP effect did not impact purge loss or steak cook loss ($P > 0.23$), but there
192 was a tendency ($P = 0.08$) for ExGP use to affect ground beef cook loss with ground beef patties
193 from the EGP- tending to have a cook loss 1.06% greater than the EGP+ treatment.

194 Warner-Bratzler shear force was measured on strip loin steaks fabricated from enhanced
195 loins (Table 3). There was a tendency ($P = 0.06$) for a dietary program \times ExGP interaction to
196 affect shear force. Dietary program influenced ($P = 0.05$) WBSF of steaks from enhanced loins.
197 The shear force of steaks from PN loins was 0.15 kg less ($P = 0.05$) than the shear force of steaks
198 from CON loins. The use of ExGP during the finishing phase did not impact ($P = 0.54$) shear
199 force of steaks fabricated from enhanced loins.

200 *3.3 Expressible moisture and water binding ability*

201 There was no dietary program \times ExGP \times period interaction for EM or WBA (Table 4; P
202 > 0.36). The two-way interactions of dietary program \times ExGP or dietary program \times period did
203 not affect EM or WBA (Table 5; $P > 0.84$). The ExGP \times period interaction did not impact ($P =$
204 0.13) EM, but did influence WBA ($P < 0.01$). Before injection, EGP+ loins had greater ($P =$
205 0.01) WBA compared to EGP- loins. After loins were injected and stored EGP- loins had a

206 greater ($P < 0.01$) WBA than EGP+ loins. Finally, the main effects of dietary program and ExGP
207 did not impact EM or WBA ($P > 0.21$).

208 **4. Discussion**

209 Beef palatability is influenced by many attributes, including tenderness, marbling,
210 texture, juiciness, and flavor profile (Miller, Carr, Ramsey, Crockett, & Hoover, 2001). These
211 attributes can be influenced by a wide range of factors, including genetics, age, diet, pre-harvest
212 management, and post-mortem processing (Hocquette, et al., 2012). Although the amount of
213 marbling often influences juiciness the most (O'Quinn et al., 2011), water holding capacity of
214 beef may also play a role in a consumer's beef eating experience (Winger & Hagyard, 1994).
215 Intrafiber water content has been suggested to impact meat tenderness (Currie & Wolfe, 1980)
216 and could potentially impact overall palatability of beef. In addition to influencing palatability,
217 poor water holding capacity of products can result in loss of salable weight for processors and
218 retailers (Huff-Lonergan and Lonergan, 2005).

219 Water holding capacity of beef and other meat products is largely influenced by pH
220 (Honikel, 2004). When meat reaches its isoelectric point the net charge of proteins is zero. This
221 effectively decreases the amount of water held between proteins as a result of the close
222 association between positively and negatively charged protein moieties. When the pH of meat is
223 transitioned away from its isoelectric point, water holding capacity improves (Aberle et al.,
224 2003). The increase in meat pH following the use of phosphates in brine solutions is well
225 documented (Smith, Simmon, McKeith, Bechtel, & Brady, 1984; Robbins et al., 2002; Wicklund
226 et al., 2005). In the present study, the common increase in pH due to brine injection was
227 duplicated. Loin pH recorded post-injection and after 7 d of storage were increased by 0.32 and

228 0.33 pH values, respectively when compared to pH recorded prior to injection. Similar to the
229 present study, Wicklund et al. (2005) reported increases in muscle pH (0.32 units) when beef
230 strip steaks were enhanced with a brine containing phosphates. Pietrasisk and Janz (2009)
231 reported a smaller pH increase of 0.23 units in enhanced beef *Semitendinosus* steaks compared to
232 non-enhanced steaks, while Robbins et al. (2002) observed that pH values of enhanced beef
233 rounds were 0.28 units greater compared to non-enhanced rounds. Finally, Grobbel, Dikeman,
234 Hunt, & Milliken (2008) reported that pH of enhanced *Longissimus*, *Semitendinosus*, and *Triceps*
235 *brachii* steaks were 0.3, 0.2, and 0.2 units higher, respectively, compared to non-enhanced
236 steaks. The current data also indicate that the ExGP \times period interaction, dietary treatment, and
237 ExGP treatment affected pH. When examining the differences closely, the differences in pH are
238 all less than 0.10, which presumably would have a relatively small biological effect on water
239 holding characteristics. Therefore, any treatment differences in water binding characteristics,
240 cook loss, WBSF post-injection and storage occurred independent of pH.

241 Water binding ability measures the capacity of muscle to assimilate and retain additional
242 water. In the current study, the WBA of EGP+ loins was greater than EGP- loins prior to
243 enhancement and storage. After enhancement, WBA of EGP+ loins were less than EGP- loins.
244 Because there were no differences in purge loss, the WBA data may indicate EGP+ were
245 incapable of retaining additional moisture after they were enhanced 110%. Pietrasik and Janz
246 (2009) found that freezing greatly reduced the WBA of meat and injection with a 1.5%
247 phosphate solution increased WBA over injection with a 0.5% phosphate solution. Since all meat
248 was frozen and the same brine solution was utilized on all loins, these factors can be eliminated
249 as contributing to the ExGP difference. It is feasible that increased WBA of EGP+ loins was due
250 to an increase in cross-sectional areas of type IIA and type IIX fibers as reported by Phelps et al.

251 (2014). Offer and Trinick (1983) stated that the majority of water present in meat is located
252 within myofibrils, and these authors demonstrated that the degree of swelling by myofibrils
253 observed after injecting brine solution was similar to the amount of water retained by the muscle.
254 Therefore, since EGP+ loins possessed greater muscle fiber cross-sectional area as reported by
255 Phelps et al. (2014), they likely possessed more myofibrils with which to bind the brine solution.

256 Previously, Phelps et al. (2014) reported that fresh loins from PN Beef Program
257 supplemented steers had less purge loss during storage and also lost less weight during cooking
258 compared to their CON counterparts. Additionally, the authors observed similar moisture
259 retention properties for loins from cattle produced without ExGPs compared to those from cattle
260 produced with ExGPs. The moisture retention advantages observed by Phelps et al. (2014) were
261 expected to be duplicated in the meat systems examined in the present study. Data in this study
262 indicate that when loins from all pre-harvest treatments were subjected to needle-injection
263 enhancement, there are no differences in purge loss or cook loss. Numerous studies indicate that
264 enhancement of fresh meat can affect both purge loss and weight loss during cooking. In a study
265 in which purge loss was compared between non-enhanced and enhanced beef and bison steaks,
266 Pietrasik, Dhanda, Shand, and Pegg (2006) reported that enhanced steaks lost 0.97% more purge
267 than non-enhanced steaks. In contrast, Pietrasik and Janz (2009) reported that enhanced
268 *Semitendinosus* steaks lost less purge than non-enhanced samples. McGee, Henry, Brooks, Ray,
269 & Morgan (2003) reported that injecting beef round roasts with a brine decreased weight loss
270 during cooking by 7.65% compared to non-injected controls, while Pietrasik and Shand (2005)
271 reported injection of *Semimembranosus* roasts decreased weight loss during cooking by 11.9%.

272 When compared to the Phelps et al. (2014) results, the lack of purge and cook loss
273 differences of enhanced loins and steaks in the current study may be a function of freezing and

274 thawing of the loins prior to enhancement. Expressible moisture, which is greatly related to
275 moisture retention during storage and cooking, was unaffected by treatment before or after
276 enhancement. This could indicate that freezing/thawing may have masked the fresh meat
277 treatment related differences in moisture retention properties that were observed by Phelps et al.
278 (2014). Boles and Swan (2002) reported that beef inside rounds that were frozen prior to
279 injection retained 1% more brine solution than fresh injected rounds. The authors concluded that
280 the disruption of the protein structure due to freezing (Polymendis, 1978) allowed for increased
281 brine uptake. Pietrasik and Janz (2009) found that when frozen/thawed meat was stored 7 d post-
282 injection, expressible moisture was drastically improved when compared to frozen/thawed meat
283 stored 1 d post injection. Since Phelps et al. (2014) did not measure expressible moisture prior to
284 freezing, it is impossible to determine the effect of freezing/thawing on this water holding
285 characteristic.

286 Enhancement of beef cuts with brine solutions has become a more common production
287 practice to alleviate tenderness issues (McGee et al., 2003). Regardless of treatment, average
288 reduction in shear force compared to Phelps et al. (2014) reported values were 13.0 N for all
289 steaks. Although it must be acknowledged the reduction in shear force of the enhanced steaks
290 may also be a function of freezing. Robbins et al. (2002) reported shear force of enhanced
291 *Semimembranosus* steaks was 0.52 kg lower than non-enhanced controls. In a study on the
292 enhancement of beef chuck muscles, Molina, Johnson, West, & Gwartney (2005) found needle
293 injection reduced WBSF of oven-roasted *Complexus* and *Subscapularis* steaks and grilled
294 *Triceps brachii* steaks compared to non-injected controls by 11.7, 12.7, and 9.9 N, respectively.
295 Also, Grobbel et al. (2008) examined the enhancement of *Longissimus lumborum*,
296 *Semitendinosus*, and *Triceps brachii* steaks and reported needle injection decreased WBSF by

297 1.73, 1.25, and 1.25 kg, respectively compared to non-injected controls. In contrast to the results
298 of Phelps et al. (2014), the data in the present study indicated that enhanced CON steaks were 1.5
299 N more tender than enhanced PN steaks. Miller et al. (2001) reported that consumers can
300 distinguish between tenderness classifications at WBSF values of 0.30 kg, suggesting that
301 differences in tenderness between the enhanced CON and PN within the present study are
302 unlikely to be detectable by consumers.

303 The greatest improvement in shear force from the Phelps et al. (2014) values due to
304 enhancement was seen in EGP+ steaks, but there were no shear force differences for pre-harvest
305 use of ExGP versus not using ExGP when loins were enhanced. Enhancement reduced the shear
306 force of EGP+ by 15.2 N compared to values reported by Phelps et al. (2014). Brooks et al.
307 (2010) reported enhancement of strip loins from cattle fed zilpaterol hydrochloride reduced shear
308 force by 0.27 kg. The greater reduction in shear force in the present study compared to Brooks et
309 al. (2010) may have been influenced by freezing and thawing of loins prior to enhancement. As
310 stated before, ice crystals formed during freezing can disrupt protein structure. Two studies
311 (Hiner, Madsen, & Hankins, 1945; Shanks, Wulf, & Maddock, 2002) report lower WBSF of
312 steaks frozen prior to cooking compared to steaks cooked fresh. The combined effects of
313 moisture enhancement and freezing may have caused a much greater reduction in shear force.

314 Since Phelps et al. (2014) detected differences in fresh steak cook loss due to dietary
315 program or ExGP use, effect of treatments on cook loss from ground beef patties was examined.
316 In contrast to the fresh steak data, there were no differences in moisture loss during cooking of
317 ground beef patties. Two mechanisms could be responsible for this finding. First, as indicated
318 throughout this paper, the cellular disruption events associated with freezing/thawing the meat
319 may have negated the increased moisture retention associated with EGP- and PN treatments

320 reported by Phelps et al. (2014). This can be inferred by the fact that all treatment groups had
321 similar expressible moisture percentages prior to pumping, therefore indicating they have similar
322 water binding capacity. The second mechanism responsible for lack of moisture retention
323 differences could be that fat was added to each sample to achieve the same final fat blend. Roth
324 et al. (1999) found that reduced-fat patties possessed more water and less fat in the raw and
325 cooked state than high-fat patties. More importantly, the authors reported that cook loss
326 percentage mirrored fat content of the blend rather than water content of the blend. This would
327 indicate that water binding is more important for ground beef blends made with a lower fat
328 content, and the differences reported by Phelps et al. (2014) may have been masked by adding
329 additional fat to the lean.

330 **5. Conclusion**

331 Injection of frozen/thawed beef loins with 0.5% sodium chloride and 0.3% sodium
332 tripolyphosphate increased loin pH by up to 0.32 units after 7-d of storage. As expected,
333 enhanced steaks had improved WBSF when compared to values reported by Phelps et al. (2014)
334 fresh steaks, but no large differences in WBSF were detected between dietary programs and use
335 of ExGP once steaks were enhanced. Expressible moisture analysis revealed that both before and
336 after injection, there was no differences in treatments to hold water. This finding is reflected in
337 the lack of pre-harvest treatment differences for purge loss and cook loss from ground beef and
338 enhanced steaks during cooking. Although not measured, freezing of the meat may have
339 influenced results in this study, thus warranting further investigation of the impact of freezing on
340 water holding capacity of beef from cattle supplemented PN Beef Program supplements.

Tables

Table 1. Diets (dry basis) for steers fed conventional feedlot diets[†] or Alltech Programmed Nutrition Beef Program[‡]

Ingredient, %	Conventional	Alltech
Wet corn gluten feed	35.00	35.00
Steam-flaked corn	53.55	53.56
Ground wheat straw	7.00	7.00
Feed additive premix	2.16	-
Mineral/vitamin supplement	2.29	2.23
Programmed Nutrition supplement	-	2.21

[†]Conventional diets included vitamin A at 2,200 IU/kg; vitamin E at 22 IU/kg; copper sulfate to provide 10 mg/kg Cu; cobalt carbonate to provide 0.15 mg/kg Co; ethylenediamine dihydriodide to provide 0.5 mg/kg I; manganous sulfate to provide 60 mg/kg Mn; sodium selenite to provide 0.3 mg/kg Se; zinc sulfate to provide 60 mg/kg Zn on a dry matter basis; as well as 300 mg•animal⁻¹•d⁻¹ of monensin and 90 mg•animal⁻¹•d⁻¹ of tylosin (Elanco Animal Health; Greenfield, IN).

[‡]The Alltech diet included Programmed Nutrition Receiver in the total mixed ration for the first 21 days at the rate of 14 g•d⁻¹•steer⁻¹ which contained: zinc proteinate to provide 10.7 mg/kg Zn; manganese proteinate to provide 7.1 mg/kg manganese; cobalt proteinate to provide 1.2 mg/kg cobalt; copper proteinate to provide 2.9 mg/kg copper; calcium iodate to provide 0.6 mg/kg iodine; selenium yeast to provide 0.31 mg/kg selenium on a dry matter basis as well as ascorbic acid, *Aspergillus oryzae* fermentation product, *Lactobacillus acidophilus* fermentation product, and *Enterococcus faecium* fermentation product. Thereafter, Programmed Nutrition Finisher was included in the total mixed ration at the rate of 20 g•d⁻¹•steer⁻¹: 10.7 mg/kg Zn; manganese proteinate to provide 7.1 mg/kg manganese; cobalt proteinate to provide 1.2 mg/kg cobalt; copper proteinate to provide 2.9 mg/kg copper; calcium iodate to provide 0.6 mg/kg iodine; selenium yeast to provide 0.31 mg/kg selenium on a dry matter basis as well as ascorbic acid, *Aspergillus niger* fermentation product, *Lactobacillus acidophilus* fermentation product, and *Enterococcus faecium* fermentation product. Both supplements were premixed into a ground corn carrier and subsequently blended into the total mixed ration.

Table 2. pH of beef strip loins from steers of steers fed conventional diets[†] or Alltech Programmed Nutrition Beef Program[‡] with and without exogenous growth promotants[§] recorded pre-injection, post-injection, and after 7 d of storage.

	Conventional		Alltech PN		SEM	<i>P</i> – value						
	EGP-	EGP+	EGP-	EGP+		Program	ExGP	Period	Prog ×Period	ExGP × Period	Prog × ExGP	Prog × ExGP × Period
pH					0.03	0.02	0.052	<0.01	0.10	0.02	0.39	0.55
Pre-injection	5.53	5.52	5.53	5.53								
Post-injection ¹	5.85	5.76	5.93	5.85								
After storage ²	5.86	5.83	5.85	5.88								

[†]Conventional diets included vitamin A at 2,200 IU/kg; vitamin E at 22 IU/kg; copper sulfate to provide 10 mg/kg Cu; cobalt carbonate to provide 0.15 mg/kg Co; ethylenediamine dihydriodide to provide 0.5 mg/kg I; manganous sulfate to provide 60 mg/kg Mn; sodium selenite to provide 0.3 mg/kg Se; zinc sulfate to provide 60 mg/kg Zn on a dry matter basis; as well as 300 mg•animal⁻¹•d⁻¹ of monensin and 90 mg•animal⁻¹•d⁻¹ daily of tylosin (Elanco Animal Health; Greenfield, IN).

[‡] The Alltech diet included Programmed Nutrition Receiver in the total mixed ration for the first 21 days at the rate of 14 g•animal⁻¹•d⁻¹ which contained zinc proteinate to provide 10.7 mg/kg Zn; manganese proteinate to provide 7.1 mg/kg manganese; cobalt proteinate to provide 1.2 mg/kg cobalt; copper proteinate to provide 2.9 mg/kg copper; calcium iodate to provide 0.6 mg/kg iodine; selenium yeast to provide 0.31 mg/kg selenium on a dry matter basis; as well as ascorbic acid, *Aspergillus oryzae* fermentation product, *Lactobacillus acidophilus* fermentation product, and *Enterococcus faecium* fermentation product. Thereafter, Programmed Nutrition Finisher was included in the total mixed ration at the rate of 20 g•animal⁻¹•d⁻¹ which contained zinc proteinate to provide 10.7 mg/kg Zn; manganese proteinate to provide 7.1 mg/kg manganese; cobalt proteinate to provide 1.2 mg/kg cobalt; copper proteinate to provide 2.9 mg/kg copper; calcium iodate to provide 0.6 mg/kg iodine; selenium yeast to provide 0.31 mg/kg selenium on a dry matter basis; as well as ascorbic acid, *Aspergillus niger* fermentation product, *Lactobacillus acidophilus* fermentation product, and *Enterococcus faecium* fermentation product. Both supplements were premixed into a ground corn carrier and subsequently blended into the total mixed ration.

[§] Exogenous growth promotants included Component ES implant administered on d 1 of the experiment, Component TE-IS at reimplant, and ractopamine hydrochloride (Optaflexx) fed at 400 mg/animal daily for the final 28 d before harvest (all products from Elanco Animal Health).

¹ Recorded following injection after a 2 min rest.

² Recorded after 7 d of storage.

Table 3. Purge loss of injected beef *Longissimus lumborum*, cook loss of injected beef *Longissimus* steaks and ground beef, and Warner-Bratzler shear force (WBSF) from steers fed conventional diets[†] or Alltech Programmed Nutrition Beef Program[‡] with and without exogenous growth promotants[§]

Item	Conventional		Alltech PN		SEM	<i>P</i> - value		
	EGP-	EGP+	EGP-	EGP+		Program	ExGP	Prog × ExGP
Purge loss ¹ , %	1.31	1.51	1.45	1.09	0.23	0.54	0.72	0.23
Steak cook loss ² , %	22.89	21.02	23.28	22.60	1.03	0.35	0.23	0.57
Ground beef cook loss ² , %	23.36	21.72	22.89	22.41	0.59	0.86	0.08	0.34
WBSF, N	20.9	19.0	18.0	19.0	1.0	0.05	0.54	0.06

[†]Conventional diets included vitamin A at 2,200 IU/kg; vitamin E at 22 IU/kg; copper sulfate to provide 10 mg/kg Cu; cobalt carbonate to provide 0.15 mg/kg Co; ethylenediamine dihydriodide to provide 0.5 mg/kg I; manganous sulfate to provide 60 mg/kg Mn; sodium selenite to provide 0.3 mg/kg Se; zinc sulfate to provide 60 mg/kg Zn on a dry matter basis; as well as 300 mg/animal daily of monensin and 90 mg/animal daily of tylosin (Elanco Animal Health; Greenfield, IN).

[‡] The Alltech diet included Programmed Nutrition Receiver in the total mixed ration for the first 21 days at the rate of 14 g/animal daily which contained zinc proteinate to provide 10.7 mg/kg Zn; manganese proteinate to provide 7.1 mg/kg manganese; cobalt proteinate to provide 1.2 mg/kg cobalt; copper proteinate to provide 2.9 mg/kg copper; calcium iodate to provide 0.6 mg/kg iodine; selenium yeast to provide 0.31 mg/kg selenium on a dry matter basis; as well as ascorbic acid, *Aspergillus oryzae* fermentation product, *Lactobacillus acidophilus* fermentation product, and *Enterococcus faecium* fermentation product. Thereafter, Programmed Nutrition Finisher was included in the total mixed ration at the rate of 20 g/animal daily which contained zinc proteinate to provide 10.7 mg/kg Zn; manganese proteinate to provide 7.1 mg/kg manganese; cobalt proteinate to provide 1.2 mg/kg cobalt; copper proteinate to provide 2.9 mg/kg copper; calcium iodate to provide 0.6 mg/kg iodine; selenium yeast to provide 0.31 mg/kg selenium on a dry matter basis; as well as ascorbic acid, *Aspergillus niger*

fermentation product, *Lactobacillus acidophilus* fermentation product, and *Enterococcus faecium* fermentation product. Both supplements were premixed into a ground corn carrier and subsequently blended into the total mixed ration.

[§] Exogenous growth promotants included Component ES implant administered on d 1 of the experiment, Component TE-IS at reimplant, and ractopamine hydrochloride (Optaflexx) fed at 400 mg/animal daily for the final 28 d before harvest (all products from Elanco Animal Health).

¹ [(pumped weight-stored weight)/pumped weight] × 100.

² [(pre-cook weight-cooked weight)/pre-cook weight] × 100.

Table 4. Expressible moisture and water binding ability of beef strip loins from steers fed conventional diets[†] or Alltech Programmed Nutrition Beef Program[‡] with and without exogenous growth promotants[§] before injection and after 7 d of storage.

Item	Conventional		Alltech PN		SEM	<i>P</i> – value						
	EGP-	EGP+	EGP-	EGP+		Program	ExGP	Period	Prog × Period	ExGP × Period	Prog × ExGP	Prog × ExGP × Period
Expressible moisture ¹ , %					0.76	0.97	0.90	0.67	0.99	0.13	0.84	0.36
Before injection	11.12	10.03	10.84	10.36								
After storage	10.20	11.36	10.75	10.85								
Water binding ability ² , %					2.05	0.96	0.21	<0.01	0.70	<0.01	0.50	0.99
Before injection	19.86	31.64	20.42	28.88								
After storage	32.79	25.61	34.21	25.49								

[†]Conventional diets included vitamin A at 2,200 IU/kg; vitamin E at 22 IU/kg; copper sulfate to provide 10 mg/kg Cu; cobalt carbonate to provide 0.15 mg/kg Co; ethylenediamine dihydriodide to provide 0.5 mg/kg I; manganous sulfate to provide 60 mg/kg Mn; sodium selenite to provide 0.3 mg/kg Se; zinc sulfate to provide 60 mg/kg Zn on a dry matter basis; as well as 300 mg•animal⁻¹•d⁻¹ of monensin and 90 mg•animal⁻¹•d⁻¹ of tylosin (Elanco Animal Health; Greenfield, IN).

[‡] The Alltech diet included Programmed Nutrition Receiver in the total mixed ration for the first 21 days at the rate of 14 g•animal⁻¹•d⁻¹ which contained zinc proteinate to provide 10.7 mg/kg Zn; manganese proteinate to provide 7.1 mg/kg manganese; cobalt proteinate to provide 1.2 mg/kg cobalt; copper proteinate to provide 2.9 mg/kg copper; calcium iodate to provide 0.6 mg/kg iodine; selenium yeast to provide 0.31 mg/kg selenium on a dry matter basis; as well as ascorbic acid, *Aspergillus oryzae* fermentation product, *Lactobacillus acidophilus* fermentation product, and *Enterococcus faecium* fermentation product. Thereafter, Programmed Nutrition Finisher was included in the total mixed ration at the rate of 20 g•animal⁻¹•d⁻¹ which contained zinc proteinate to provide 10.7 mg/kg Zn; manganese proteinate to provide 7.1 mg/kg manganese; cobalt proteinate to provide 1.2 mg/kg cobalt; copper proteinate to provide 2.9 mg/kg copper; calcium iodate to provide 0.6 mg/kg iodine; selenium yeast to provide 0.31 mg/kg selenium on a dry matter basis; as well as ascorbic acid, *Aspergillus niger* fermentation product, *Lactobacillus acidophilus* fermentation product, and *Enterococcus faecium* fermentation product. Both supplements were premixed into a ground corn carrier and subsequently blended into the total mixed ration.

[§] Exogenous growth promotants included Component ES implant administered on d 1 of the experiment, Component TE-IS at reimplant, and ractopamine hydrochloride (Optaflexx) fed at 400 mg/animal daily for the final 28 d before harvest (all products from Elanco Animal Health).

¹ [(Initial weight-centrifuged weight)/initial weight] × 100.

² Calculated as 300-(11.43 × supernatant volume).

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