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9 **Physical and Biochemical Mechanisms Associated with Beef Carcass Vascular Rinsing**

10 **Effects on Meat Quality: A Review**

11

12 **Abstract**

13 Carcass vascular rinsing and chilling involves infusing a chilled isotonic solution (98.5% water  
14 and a blend of mono- and di-saccharides and phosphates) into the vasculature immediately upon  
15 exsanguination. Primary purposes of carcass vascular rinsing are to (1) effectively remove residual  
16 blood from the carcass; (2) lower internal muscle temperature rapidly; and (3) optimize pH decline  
17 by effective delivery of glycolytic substrates in the rinse solution. Previous studies have revealed  
18 that the beef carcass vascular rinsing early postmortem positively affects meat quality, product  
19 shelf-life, and food safety. Thus, the objective of this review is to provide a more comprehensive  
20 understanding of the physical and biochemical mechanisms associated with beef carcass vascular  
21 rinsing, focusing on the relationship between quality attributes (CIE L\*a\*b\*, chemical states of  
22 myoglobin, oxygen consumption and sarcomere length) and muscle metabolic response to various  
23 substrate solutions (Rinse & Chill<sup>®</sup>, fructose, sodium phosphate, and dipotassium phosphate) that  
24 stimulate or inhibit the rate of glycolysis early postmortem. In addition, this review discusses the  
25 absence of metabolite residues (phosphorus, sodium, and glucose) related to the application of the  
26 chilled isotonic solution. This review primarily focuses on beef and as such extending the  
27 understanding of the mechanisms and meat quality effects discussed to other species associated  
28 with vascular rinsing, in particular pork, may be limited.

29

30 **Key Words** – beef, carcass chilling, anaerobic glycolysis, meat quality, tenderness

31 **Introduction**

32           The novel postmortem process referred to as Rinse & Chill<sup>®</sup> technology (RC: MPSC Inc.,  
33 Hudson, Wisconsin, USA) entails inserting a sanitized catheter into the carotid artery of an animal  
34 immediately upon exsanguination, and a chilled isotonic solution is then infused into the  
35 vasculature at a rate up to 10% of the carcass weight. The RC solution pushes the blood out of the  
36 carcass through the venous vasculature (jugular veins) and also continues to drain from the carcass  
37 similar to normal bleeding (Kethavath et al., 2022; Mickelson & Claus, 2020). In detail, on the kill  
38 floor, each carcass is weighed by an automated process control system and the amount of rinse  
39 solution needed is calculated. This process requires typically approximately 4 min or less per beef  
40 carcass, and the catheter is then removed. The suspended carcass passes through the normal  
41 slaughter procedure along connecting rails.

42           The RC solution is primarily composed of water (98.5%) with a blend of dextrose, maltose,  
43 and sodium phosphates. All of the ingredients in the RC solution are approved by the U.S. Food  
44 & Drug Administration and are internationally GRAS-listed, common food-grade ingredients. The  
45 solution is designed based on the hypothesis: dextrose (glucose) is the normal substrate in muscle  
46 used in muscle metabolism to produce energy. Maltose is simply a disaccharide composed of two  
47 glucose units which the muscle utilizes to provide additional glucose for metabolism. Phosphates  
48 stimulate energy metabolism and are naturally present in the muscle. Phosphatases also present in  
49 the muscle rapidly hydrolyze the phosphates as part of normal muscle metabolism (Sickler et al.,  
50 2013; Kilic et al., 2019). Thus, these substrates are used to enhance the glycolytic metabolism of  
51 the muscle (Hunt et al., 2003; Yancey et al., 2001). These substrates leave no detectable residues  
52 in beef (Hwang et al., 2020) since the muscle is physiologically active at the time of vascular  
53 rinsing early postmortem.

54 Recently, a number of studies have been investigated (Hwang et al., 2020; Kethavath et al.,  
55 2020; Kethavath et al., 2022; Moreira et al., 2018) to address the physical and biochemical  
56 properties related to carcass vascular rinsing effects on beef meat quality. Therefore, this review  
57 is to provide a review of the potential physical and biochemical mechanisms associated with beef  
58 carcass vascular rinsing that impact color stability, oxygen consumption, sarcomere length,  
59 metabolite residues, and muscle contractile responses.

## 61 **Effects of beef carcass vascular rinsing on meat quality**

### 62 *Color stability*

63 Previously published studies have found that the carcass vascular rinsing has a beneficial  
64 effect on meat color stability (Hunt et al., 2003; Fowler et al., 2017; Kethavath et al., 2021 and  
65 2022; Mickelson & Claus, 2020). Common results were more red (higher CIE a\* and  
66 deoxymyoglobin, lower metmyoglobin) and lighter (higher CIE L\*) of the meat color in the *triceps*  
67 *brachii* (shoulder), *longissimus lumborum* (loin), and *semimembranosus* (ham) from a variety of  
68 animal species (beef, bison, pork, and lamb). In a study by Moreira et al. (2018), they observed  
69 RC lean ground beef resulted in greater redness (CIE a\*: 15.75 vs 13.06; P<0.05), higher  
70 deoxymyoglobin (DMb: 1.29 vs 1.12; P<0.05), and lower metmyoglobin (MMb: 0.94 vs 1.11;  
71 P<0.05) on 7 d display than the beef samples from the non-rinsed carcasses (Table 1). RC appeared  
72 lighter and more yellow on 1 d display, but no other differences were seen on day 4 and 7 between  
73 the control (CN) and RC. Associated with the increase in lightness, Farouk and Price (1994)  
74 suggested that the higher yellowness (CIE b\*) determined might be responsible for the increased  
75 lightness, as a result of the greater light scattering. Erazo-Castrejón et al. (2019) found that the RC  
76 removed 40% more residual blood from the pork muscle compared to the conventional chilled

77 carcasses. RC had lower hemoglobin (Hb, 13.6% vs 19.1%) and higher myoglobin (Hb, 86.4% vs  
78 80.9%) than the CN when the percentages of Mb and Hb by weight (relative to each other) were  
79 calculated. Thus, the additional blood removal from the carcass might have contributed to the  
80 lighter colored meat.

81 Myoglobin oxidation is one of the major non-microbial factors which result from lipid  
82 oxidation and induce quality deterioration in muscle foods. RC tended to slow down the myoglobin  
83 oxidation process with the greater DMb and lower MMb compared to CN during the display  
84 periods (Table 1). These results could be related to sodium phosphates in the RC solution that act  
85 as antioxidants and inhibit lipid oxidation by chelating metal ions ( $\text{Fe}^{2+}$ ,  $\text{Cu}^{2+}$ , etc.). Wu et al. (2021)  
86 reported that the reduced Mb forms (DMb) have lower pro-oxidant ability in muscle foods when  
87 compared to their oxidized form (MMb). In addition, Hb and Mb are the most abundant heme  
88 proteins and contribute to accelerating lipid oxidation in postmortem muscles. With more blood  
89 being removed by RC, besides the additional amount of hemoglobin that is reduced in the  
90 vasculature, RC would also remove non-heme iron. Non-heme iron has the potential to cause  
91 oxidation of lipid and myoglobin. This is also closely associated with the flavor which is negatively  
92 affected by the hemoglobin and non-heme iron. Yancey et al. (2002) reported that steaks  
93 (*semitendinosus*) from the RC beef carcasses had a lower cardboard flavor (rancidity) compared  
94 to the non-rinsed samples. A greater beef flavor was identified in cooked ground beef from the RC  
95 cattle than the non-rinsed carcasses. Thus, the enhanced blood removal and sodium phosphates  
96 likely contributed to the meat redness, an extension in meat color stability, and the sensory quality.

97

98 *Oxygen consumption*

99 Oxygen consumption represents the ability of postmortem muscle to consume oxygen. In  
100 postmortem muscle, there is competition for oxygen among mitochondria, oxygen-consuming  
101 enzymes, myoglobin, microorganisms, and lipid and protein oxidation (Ramanathan et al., 2017).  
102 In particular, the competition between myoglobin and mitochondria is the key factor that  
103 influences the formation of bright cherry red color (oxymyoglobin). If mitochondrial activity  
104 outcompetes myoglobin for oxygen, this will result in a darker, deoxygenated meat color  
105 (predominant DMb) due to limited oxygen supply to myoglobin (Mancini, 2009). In addition, if  
106 the oxygen level in the meat is very low, this promotes MMb formation.

107 RC ground beef tended to have a greater oxygen consumption as the amount of residual  
108 oxygen was lower than CN, but no difference was observed in the ground loin (Table 2; Kethavath  
109 et al., 2020). However, Kethavath et al. (2021) found that RC had greater oxygen consumption  
110 (RC 4.56 vs CN 5.18 % O<sub>2</sub>; P<0.05) immediately after removing the PVC film and vacuum  
111 packaging of the ground pork shoulder. Other studies that just compared muscles (non-rinsed  
112 muscles) described that the psoas muscle had greater oxygen consumption than the longissimus  
113 muscle, suggesting that this muscle predominantly consisted of red fibers, and accordingly greater  
114 oxidative metabolism was than the longissimus (Mohan et al., 2010; Ke et al., 2017). In another  
115 study (Ramanathan and Mancini, 2018), greater oxygen consumption can contribute to enhanced  
116 color stability. This likely involves mitochondrial production of NADH and other reducing  
117 equivalents related to maintaining myoglobin in ferrous form that is associated with the formation  
118 of deoxymyoglobin and oxymyoglobin.

119 RC promptly removes more oxygen from oxygen-carrying hemoglobin by the enhanced  
120 blood removal that stimulates the rate of postmortem anaerobic glycolysis. Perhaps this shift from  
121 aerobic metabolims to anaerobic metabolism facilitates preserving mitochondria activity reflected

122 in the greater oxygen consumption in the RC muscles. The aerobic metabolism through the TCA  
123 cycle induces production of more NADH that helps maintain the heme iron in the ferrous state  
124 (Kethavath et al., 2020). With greater oxygen consumption and generation of NADH would  
125 facilitate formation of deoxymyoglobin which is a more stable oxidative state than oxymyoglobin.  
126 Therefore, RC has the potential to positively affect color development and color stability (limit  
127 oxidation) in beef that is anaerobically packaged.

128

### 129 *Sarcomere length*

130 When muscles enter rigor mortis, sarcomere shortening can occur, depending on chilling  
131 conditions (e.g., rate, temperature), carcass suspension methods, and glycolytic metabolism within  
132 the muscle fibers that will have effects on postmortem energy metabolism, pH development, and  
133 proteolytic activity (Warner et al., 2014). It is well established that when a carcass temperature  
134 declines below 10°C early postmortem while the pH is still above 6 and the presence of adenosine  
135 5' triphosphate (ATP), the muscle can shorten, caused by excessive release of calcium ions from  
136 the sarcoplasmic reticulum (Davey et al., 1976; Tornberg, 1996). In contrast, a rapid drop in pH  
137 while a carcass temperature is still warm (35–42°C) can lead to negative effects on color (pigment  
138 denaturation) and reduction in water holding capacity due to the less physical space for water to  
139 bind to the contractile proteins of actin and myosin (Hopkins et al., 2014, Mickelson & Claus,  
140 2020; Warner et al., 2021).

141 Kethavath et al. (2022) conducted research with beef loins and determined the sarcomere  
142 length, wherein sarcomeres in the beef loin of RC were approximately 27% longer (Table 2) than  
143 in the CN. They noted that electrical stimulation was not applied to the CN or RC carcasses to  
144 accelerate rigor mortis and help avoid cold shortening. In the absence of electrical stimulation and



145 since the muscles were excised from the carcass at 24 h postmortem, perhaps rigor was not yet  
146 completed, and the muscles were capable of continuing to shorten. In contrast to the control,  
147 Kethavath et al. (2022) found that RC accelerated the rate of anaerobic glycolysis and the pH  
148 declined before conditions known to cause cold-induced occurred. In RC the pH was below 6 when  
149 the temperature reached approximately 10-15°C (>12 h postmortem) whereas in the control the  
150 pH was 6.3. This finding confirms those by Monin and Santé-Lhoutellier (2014) who suggested  
151 that the ideal temperature for meat to prevent cold shortening is approximately 15°C at rigor mortis  
152 (>12 h postmortem without stimulation).

153 A potential alternative hypothesis maybe related to RC stimulating early release of calcium  
154 from the sarcoplasmic reticulum and limiting the ability of this membrane system to adequately  
155 resequenter calcium (Mickelson and Claus, 2020). When carcasses are vascularly rinsed, the  
156 appendages typically extend and stiffen. This physical change may be related to the released  
157 calcium or perhaps the physical effect of the pressure of the rinse solution during the application.  
158 After the vascular rinsing terminates the appendages mostly returned to their pre-rinsed anatomical  
159 position. Once the muscle has limited ATP, the sarcoplasmic reticulum would not be able to  
160 resequenter the calcium and the muscle would be locked into rigor thereby the muscle would be  
161 unaffected by temperatures below 10 °C. Another aspect of RC is that the early release of calcium  
162 could encourage more desirable conditions for calpain activity and affect the contractile  
163 mechanism. Calpains are known to increase in activity when calcium is available and cause  
164 tenderization. Several studies related to the beef carcass vascular rinsing have shown improvement  
165 in tenderness, wherein tenderness was improved by 20% in cow striploin steaks (Hite et al., 2019),  
166 24% in bison steaks (Mickelson & Claus, 2020), 56% in steaks from light dairy cows, and 58% in  
167 steaks from lean dairy cows (Kethavath et al., 2022). Thus, the chilled isotonic RC solution appears

168 to counterbalance potential unfavorable changes related to a more rapid pH decline, as evidenced  
169 by greater sarcomere length in the beef loin. In addition, despite the more rapid pH decline in the  
170 carcass, use of the chilled RC solution and its effect on efficiently removing heat out of the carcass  
171 helps protect the meat pigments from being denatured and improves the red color stability.

172

### 173 *Muscle contraction*

174 Contractile system of the muscle can be stimulated depending on the release of calcium.  
175 Generally, muscle contractions are accelerated as postmortem metabolism proceeds under  
176 anaerobic conditions. As a result, lactate and hydrogen ions which are the end products of  
177 anaerobic glycolysis are accumulated in the muscle (Matarneh et al., 2017). The muscle pH  
178 normally lowers from approximately 7.2 in living muscle to an ultimate pH between 5.5 and 5.7.  
179 The rate and extent of postmortem metabolism is a critical factor affecting protein functionality  
180 and meat quality attributes such as color, texture, flavor, water holding capacity, and shelf-life  
181 (Hopkins et al., 2014, Warner et al., 2021). However, in spite of its importance, the rate and extent  
182 of muscle metabolism and its role during postmortem glycolysis have not been clearly established  
183 associated with vascular rinsing effects.

184 Determination of contractile force responses in pre-rigor muscles were studied to  
185 understand if metabolic activity could be modulated by various glycolytic substrates. Recently, a  
186 preliminary study was done by Moreira et al. (2019) who determined muscle contractile responses  
187 to electrical stimulation from the beef *sternomandibularis* muscles exposed to various substrate  
188 solutions (Figure 1). It was demonstrated that the contraction force decreased as time increased.  
189 Faster declines in contraction force were observed using the standard Rinse & Chill® solution as  
190 well as fructose, whereas the forces were maintained longer and decreased slowly with the use of

191 dipotassium phosphate and sodium phosphate. Regarding the faster decline in force, this is likely  
192 due to the increased rate of postmortem glycolysis. The accelerated release of calcium then  
193 contributes to a rapid muscle contraction and an increase in the rate of muscle metabolism and the  
194 rate of pH decline (Huff-Lonergan and Lonergan, 2005). Besides, the cold temperature of the  
195 solution likely affects the ability of the terminal cisternae of the sarcoplasmic reticulum to store  
196 calcium when carcasses are vascularly rinsed. Matarneh et al. (2017) stated that at lower  
197 temperatures, the sarcoplasmic reticulum actively pumps calcium ions out of the sarcoplasm while  
198 calcium sequestering is impaired, thereby promoting the cytosolic concentration of calcium. Thus,  
199 this result confirms RC leads to early release of calcium ions from the sarcoplasmic reticulum,  
200 having a direct effect on activation of the muscle shortening (Devine et al., 2014).

201

#### 202 *Metabolite residues*

203 A study has revealed that chilling method did not impact ( $P>0.05$ ) metabolite residues  
204 (Table 3; phosphorus, sodium, and glucose) in the ground beef loin (Hwang et al., 2020). They  
205 also reported that their results were similar to numerous studies (Czerwonka et al., 2015; Flowers  
206 et al. 2018; Garmyn et al., 2011; Mateescu et al., 2013), wherein phosphorus and sodium were  
207 naturally found in loin muscles from conventionally chilled beef carcasses. In detail, the results of  
208 phosphorus were: CN = 1,667  $\mu\text{g/g}$ ; RC = 1,661  $\mu\text{g/g}$ . The concentrations of phosphorus naturally  
209 found in beef were 1,727  $\mu\text{g/g}$  in steer, 1,945  $\mu\text{g/g}$  in cow, 2,167  $\mu\text{g/g}$  in bull, and 2,022  $\mu\text{g/g}$  in  
210 cattle, respectively. The sodium content was 711  $\mu\text{g/g}$  in CN and 655  $\mu\text{g/g}$  for RC. The levels of  
211 sodium naturally present in beef were 352  $\mu\text{g/g}$  in steer, 530  $\mu\text{g/g}$  in cow, 510  $\mu\text{g/g}$  in cattle, and  
212 533  $\mu\text{g/g}$  in bull, respectively. The glucose contents (CN 6.81  $\mu\text{mol/g}$  vs RC 7.49  $\mu\text{mol/g}$ ) were  
213 very similar to those reported by Rhoades et al. (2005) in the beef *M. Sternocephalicus* pars

214 mandibularis which contained 6.54  $\mu\text{mol/g}$ . Antonelo et al (2020) found that the glucose was 4.11  
215  $\mu\text{mol/g}$  in the beef loin. Falowo (2021) noted that there are large variations in mineral content of  
216 muscle foods which are mainly affected by pre-mortem (nutrition, species, breed, sex, age at  
217 slaughter, muscle types, etc.) and post-mortem (processing methods, methods for the  
218 determination of minerals) factors.

219 Although the RC solution used to rinse out the blood from the vasculature contains some  
220 substrates. At the time of early postmortem and vascular rinsing, pre-rigor muscle is still  
221 physiologically active. These substrates are metabolized by the muscle, leaving no detectable  
222 residues compared to meat from non-rinsed carcasses. The carcasses are vascularly rinsed at no  
223 more than 10% of the carcass weight with a cold isotonic solution, and the solution is allowed to  
224 freely drain. Therefore, these findings confirmed that after postmortem storage, no differences in  
225 glucose and phosphorus residuals between the non-rinsed beef and the RC beef exist. Based on the  
226 inherent amount in sodium naturally found in beef, and the rinse solution is allowed to drain, the  
227 minor contribution associated with the phosphates does not result in a difference in the sodium  
228 content of the beef.

229

## 230 **Conclusion**

231 The primary focus of this review was on beef associated with vascularly rinsing carcasses. This  
232 review has provided a more comprehensive understanding of the potential biochemical  
233 mechanisms on how vascularly rinsing, Rinse & Chill<sup>®</sup>, modulates glycolytic activity and its  
234 effects on various meat quality outcomes. Results based on early published research combined  
235 with the recent studies revealed that Rinse & Chill<sup>®</sup> has the ability to stimulate the rate of  
236 glycolysis early postmortem that facilitates the decline in pH. Beef carcasses vascularly rinsed

237 have improved color stability and tenderness, and leave no detectable rinse solution saccharides or  
238 phosphate residues in the meat. Future research that explores an understanding of the effects of  
239 this process on sarcoplasmic reticulum functionality would be beneficial as well as a deeper  
240 understanding of how the impact on mitochondrial activity. In addition, an understanding of the  
241 effects of differences in the glycolytic status among different beef animals at the time of harvest  
242 would contribute to the understanding of the process that could lead to further optimization of  
243 desired meat quality outcomes.

244

## 245 **Acknowledgements**

246

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336 Table 1. Effects of carcass vascular rinsing<sup>1</sup> on color parameters (CIE L\*, a\*, b\* and chemical  
 337 state of myoglobin) under continuous lighting display conditions on ground beef loin.

TRT	Day			Day			Day		
	1	4	7	1	4	7	1	4	7
	<b><u>Redness (CIE a*)</u></b>			<b><u>Lightness (CIE L*)</u></b>			<b><u>Yellowness (CIE b*)</u></b>		
CN	20.01 <sup>a</sup>	15.85 <sup>b</sup>	13.06 <sup>c</sup>	44.88 <sup>abc</sup>	44.35 <sup>bc</sup>	43.85 <sup>c</sup>	9.90 <sup>ab</sup>	8.70 <sup>bc</sup>	7.94 <sup>c</sup>
RC	21.65 <sup>a</sup>	17.05 <sup>b</sup>	15.75 <sup>b</sup>	46.67 <sup>a</sup>	45.69 <sup>ab</sup>	43.43 <sup>c</sup>	10.84 <sup>a</sup>	9.56 <sup>ab</sup>	7.46 <sup>c</sup>
	<b><u>Oxymyoglobin (OMb)</u></b>			<b><u>Deoxymyoglobin (DMb)</u></b>			<b><u>Metmyoglobin (MMb)</u></b>		
CN	2.18 <sup>a</sup>	1.95 <sup>b</sup>	1.76 <sup>c</sup>	1.06 <sup>b</sup>	1.06 <sup>b</sup>	1.12 <sup>b</sup>	0.87 <sup>cd</sup>	0.98 <sup>b</sup>	1.11 <sup>a</sup>
RC	2.29 <sup>a</sup>	1.96 <sup>b</sup>	1.82 <sup>c</sup>	1.06 <sup>b</sup>	1.06 <sup>b</sup>	1.29 <sup>a</sup>	0.84 <sup>d</sup>	0.97 <sup>bc</sup>	0.94 <sup>bcd</sup>

338 Source: data from Moreira et al. (2018). Larger numbers indicate greater OMb, DMb, and MMb.

339 <sup>1</sup>Carcass chilling treatment: CN, not vascularly rinsed; RC, Rinse & Chill<sup>®</sup>.

340 <sup>a-d</sup>Means within a dependent variable with unlike superscript letters are different (P<0.05).

341 Table 2. Effects of carcass vascular rinsing<sup>1</sup> on oxygen consumption and sarcomere length on beef  
342 loin.

<b>TRT</b>	<b>Oxygen Consumption (%)</b>	<b>Sarcomere Length (<math>\mu\text{m}</math>)</b>
CN	2.58 <sup>a</sup>	1.42 <sup>a</sup>
RC	2.45 <sup>a</sup>	1.80 <sup>b</sup>

343  
344 Source: oxygen consumption on ground beef loin from Kethavath et al. (2020); sarcomere length  
345 on beef loin muscle from Kethavath et al. (2022).

346 <sup>1</sup>Carcass chilling treatment: CN, not vascularly rinsed; RC, Rinse & Chill<sup>®</sup>.

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347 Table 3. Effects of carcass vascular rinsing<sup>1</sup> on residual phosphorus, sodium and glucose contents  
348 on ground beef loin.

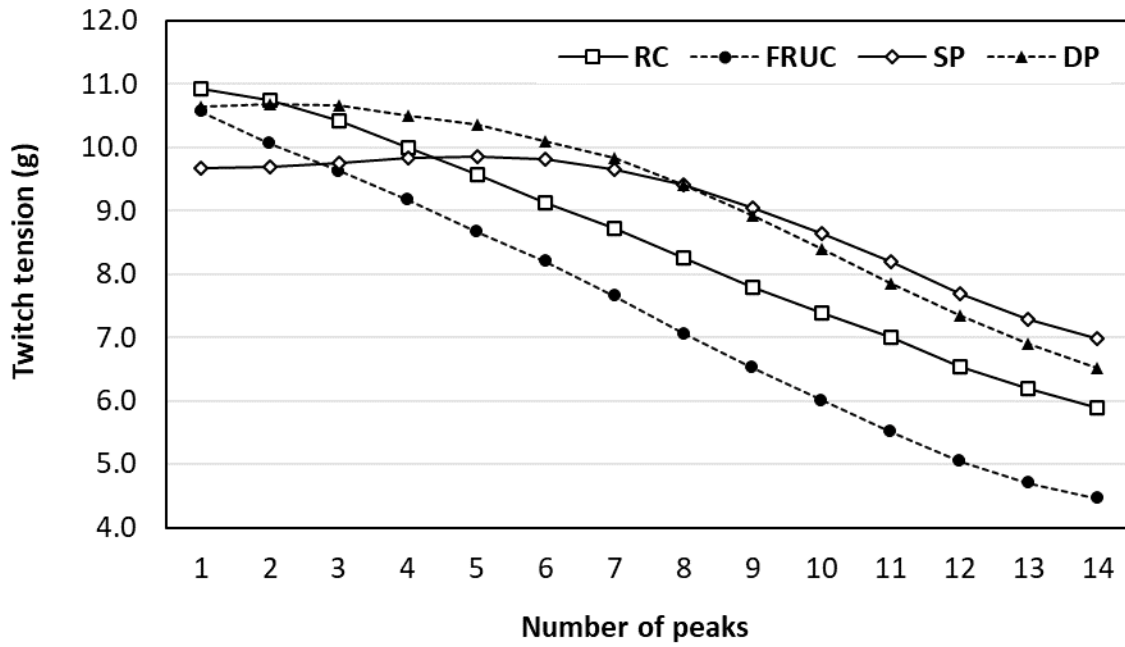
	<b>TRT</b>	<b>Phosphorus (µg/g)</b>	<b>Sodium (µg/g)</b>	<b>Glucose (µmol/g)</b>
349	CN	1666.91 <sup>a</sup>	710.77 <sup>a</sup>	6.81 <sup>a</sup>
350	RC	1661.46 <sup>a</sup>	654.58 <sup>a</sup>	7.49 <sup>a</sup>
	S.E.	77.74	30.33	0.47

351 Source: data from Hwang et al. (2020).

352 <sup>1</sup>Carcass chilling treatment: CN, not vascularly rinsed; RC, Rinse & Chill<sup>®</sup>.

353 <sup>a</sup>Means within a column are not different ( $P > 0.05$ ).

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355 Figure 1. Contractile properties of myofiber muscle exposed to four different solutions (adapted

356 from Moreira et al., 2019). RC, Rinse & Chill®; FRUC, fructose; SP, sodium phosphate; DP,

357 dipotassium phosphate.