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8	Evaluation of Meat from Korean Native chickens: Analysis of Biochemical
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Abstract

This study examined biochemical components, fatty acids, antioxidant dipeptides, and 35 muscle fiber density of breast and thigh muscles from Korean new native chicken strains (A 36 37 and B) at two slaughter ages, compared with white semi-broiler (W) or broilers. The pH values were different by chicken breed. The new native strains had the lowest fat content in the breast 38 at 12 wk (p < 0.05). Regardless of the muscles, A and B at 12 wk had higher levels of 39 arachidonic acid (ARA; C20:4), docosahexaenoic acid (DHA; C22:6), and nervonic acid 40 (C24:1) than broilers (p < 0.05). A similar result was observed for the polyunsaturated fatty 41 acids (PUFAs) and polyunsaturated and saturated fatty acids ratio (P/S) content in the breast. 42 Irrespective of the muscles, A and B enriched with omega-3 fatty acids had a lower ω -6/ ω -3 43 44 PUFA ratio than broilers (p < 0.05) at 12 wk. Of the antioxidant di-peptides, the anserine contents were highest in A and B than in the W or broilers (p < 0.05), regardless of the muscles 45 and slaughter ages. Furthermore, the breast meat from A and B contained a higher muscle fiber 46 density for both slaughter ages than the W and broilers (p < 0.05). Based on these findings, even 47 if the commercial birds (broilers or W) are raised under the similar environmental conditions 48 49 as A and B, the new native chicken strains have distinct meat quality attributes, particularly higher ARA and DHA levels, lower ω -6/ ω -3 PUFA ratio, and higher anserine contents. 50

51 Keywords native chicken, fatty acids, antioxidant dipeptides, muscle fiber density, FE-SEM

52 Introduction

White meat, such as chicken meat, is considered healthier than red meat because of its relatively lower fat, cholesterol, and iron contents (Jaturasitha et al., 2008). Unlike the global chicken meat consumption, the per capita consumption of poultry meat in Korea has risen by approximately 3 kg over the last few decades, based on a study by the Korean Ministry of Agriculture and Forestry. Globally, however, this hastily growing consumption of chicken meat is centered on a limited number of fast-growing broiler strains regulated by commercial breeding businesses in intensive fattening systems (Jaturasitha et al., 2008). On the other hand, a special attention in slow-growing chickens has been increasing recently since of the consumers' demands for healthier food (Fanatico et al., 2007).

Despite the increasing circumstances of slow-growing chickens, Korean native chickens 62 have not been raised in sufficient numbers because of their low productivity and undesirable 63 meat characteristics. Usually, native chicken meat in diverse countries has a unique taste and 64 texture that draws the attention of domestic consumers, thereby it leads the price by 65 approximately 2~3 folds that of commercial broilers (Ding et al., 1999). On the other hand, 66 however, meat quality is affected significantly by numerous factors, such as genetics, age, body 67 weight, feed, and other environmental conditions (Chen et al., 2002; Jaturasitha et al., 2008; 68 Moore et al., 2008; Mora et al., 2008). The meat pH, fat, cholesterol, fatty acids, antioxidant 69 dipeptides (carnosine and anserine), and fiber size contribute to the processing functionality 70 and consumer acceptance of meat (Jeon et al., 2010). Owing to critical characteristic for 71 successful product formulation and process control, pH has a great effect on meat quality 72 (Bianchi et al., 2005). However, the pH of meat is regulated by a wide verities of factors, such 73 as genetics (breed lines), gender, the manner of holding animals, transport, lairage conditions 74 and time - pre-slaughter stress, method of slaughter, technological parameters, postmortem 75 76 handling, and storage time of meat (Ristic and Damme, 2013). Although the genotype has a minor impact on the fatty acid profile of muscle lipids compared to feeding, a difference in 77 growth intensity caused by the genotype could still affect the fatty acids essential to human 78 79 health (Jaturasitha et al., 2008). The lipid composition of meat is influenced primarily by the birds' diet, even though other factors, such as age and breed line, also influence the fatty acid 80 profile (Popova et al., 2016). One of the most significant quality indices in chicken meat is the 81

82 cholesterol content, which needs to be as low as possible (Osada et al., 2006). Poultry strains and ingredients of the feed are the most contributing factors to decreasing the cholesterol level 83 in the meat. In addition, dietary cholesterol is strongly allied with coronary heart disease and 84 85 arteriosclerosis (Simopoulos, 2004). Recently, many challenges have been implemented to reduce the cholesterol contents in chicken meat by addition with dietary supplements, such as 86 omega-3, garlic, and copper (Chowdhury et al., 2002). However, in Korea, several researchers 87 have conducted experiments on the quality characteristics of Korean native chicken meat over 88 the past few years and showed that Korean native chicken (KNC) has nutritional qualities, 89 including a unique flavor and texture (Ali et al., 2019; Jayasena et al., 2013; Jeon et al., 2010). 90 As endogenous compound, the functional ingredients in chicken meat, such as carnosine, 91 anserine, and functional compounds, has increased considerably with a great interest (Mora et 92 93 al., 2008; Peiretti et al., 2012; Purchas et al., 2004). A recent study by Ali et al. (2019) compared the carnosine, anserine, and contents of raw meat from three different new native 94 chicken strains in Korea. It has been reported that carnosine (β -alanyl histidine) and anserine 95 96 (1-methyl carnosine), were assumed a stratagem for poultry production (Schmid, 2009). Anserine is an *N*-methylated derivative of carnosine, dominated in non-mammalian species, 97 such as poultry and it occupies the similar biological activities to carnosine (pH buffer in 98 muscles, antiglycation, antiaging, antioxidation, and neurotransmitter functions) (Schmid, 99 100 2009). Compare with carnosine, anserine is the most ample dipeptide initiated in most types of 101 poultry meat, including Korean native chickens (Ali et al., 2019; Jung et al., 2013).

The muscle fiber number and the cross-sectional area also increased with the body weight and age of poultry (Berri et al., 2007). On the other hand, the chick weight at hatching increases with increasing hen age. The morphology of the muscle arrangements in a state of original or changed after treatments can be related to eating tenderness in meat industry (Jones et al., 1977). Muscle structure differs according to the muscle types, species, and breed of animal or bird, all 107 of which contribute to the metamorphoses in muscle texture.

In Korea, the Ministry of Agriculture, Food and Rural Affairs, and Ministry of Oceans 108 and Fisheries, RDA, has conducted a native chicken breed restoration program through the 109 Golden Seed Project to maintain national resources and consumers' demand. As a part of this 110 program, a few commercial meat-type native chicken strains were developed. The different 111 quality characteristics were studied recently and compared with those of commercial broilers 112 and a commercial native chickens (Ali et al., 2019). Of them, two new native chicken strains 113 (A and B) were proposed as candidates for selection, considering their nutritional and bioactive 114 attributes under this study. Therefore, a study of the differences in the quality characteristics 115 between white semi-broiler/broiler and two native chicken strains (A, and B) under controlled 116 environmental factors is warranted. Accordingly, this study compared the quality attributes of 117 118 breast and thigh meat from white semi-broilers (W) and broilers with two new native chicken strains (A and B) raised and slaughtered at similar live weights at two production stages. 119

120

121 Materials and Methods

122 **Bird**

At an experimental farm of Harim Corporation (Gimje, Korea), 200 male chicks from new 123 native chicken strains (A and B) were reared for 5 and 12 wk respectively under similar 124 conditions and diets. Mating combinations from the paternal line and two separate maternal 125 126 lines were used to produce the new native chicken strains. W was raised for 5 wk and broiler chickens were also raised to the same slaughter weight (about 1.9 kg) as the new native chicken 127 strains at 12 wk, and used as the controls. Newly bred A and B were slaughtered alongside the 128 129 W and broiler chickens of equal live weight. The carcasses were vacuum-packed and stored at -20°C before the study. After chilling at 4°C for 24 h, the left and right sides of the carcasses' 130 breast (*pectoralis major*) and thigh (*biceps femoris*) muscles were dissected. After sorting the 131

samples, they were minced in a food mixer (Kitchen Aid, 5KPMS, USA). The goal supernatant
was then rendered from the minced meat samples under various conditions to conduct the study.

134 **Reagents and materials**

Sigma-Aldrich provided the standards (carnosine and anserine) (St. Louis, MO, USA).
Cholesterol (Tokyo Chemical Industry, Tokyo, Japan) and 5-cholestane were used as the
external and internal criteria (IS) in cholesterol analysis (Sigma-Aldrich Co., St. Louis, MO, USA).
USA).

139 Laboratory analysis

The pH in the breast and thigh from W, broiler, A, and B treatments were measured using 140 a pH meter (Seven Excellence[™], METTLER TOLEDO, Greifensee, Switzerland). For fat 141 contents, the lipids were extracted from 5 g of muscle with chloroform/methanol (2:1), 142 according to the (Folch and Lees, 1951). The carnosine and anserine contents were determined 143 using a slight modification of the method reported by Mora et al. (2007), where 2.5 g of 144 145 pulverized meat samples from each chicken were homogenized with 7.5 mL of 0.01N HCl at 1,130×g for 30 s (T25 basic, IKA GmbH & Co. KG, Germany). After homogenization, the 146 samples were centrifuged for 30 min at 4°C at 10,000 g (1580 R, GYROZEN Co., Ltd, Korea). 147 A 0.5 mL sample of centrifuged supernatant was combined with 1.5 mL of acetonitrile in a 2 148 mL tube and centrifuged for 10 min at 10,000 g at 4°C (1580 R, GYROZEN Co., Ltd). The 149 supernatant was analyzed by high-performance liquid chromatography (HPLC, 1580R, 150 GYROZEN Co., Lt) after being filtered through a 0.2 m membrane filter. 151

152 Gas chromatography (GC) was used to assess the cholesterol content of minced samples 153 (HP 5890 Series II, Hewlett Packard Co., Palo Alto, CA, USA). HP-5 ($30 \text{ m} \times 0.320 \text{ mm}$, 0.25 154 µm, Agilent, J & W Scientific, Santa Clara, CA, USA) was used as the analytical panel. The 155 fatty acids composition of the breast and thigh meat was estimated using the method reported by O'fallon et al. (2007) with a minor modification. The assay was carried out using an Agilent 7890 series Gas Chromatograph-Flame Ionization Detector under the following conditions: injector split mode with a split ratio of 25:1, and temperature 250°C. High purity air, high purity H₂, and high purity He were used as carrier gases. The flow rate for H₂ and air was 40 mL/min and 400 mL/min, respectively. Each peaks were identified using 37-FAME standard (Agilent), and a fatty acid was represented as a relative proportion of the total identified fatty acids.

163

164 Muscle fiber density

The muscle fiber characteristics were determined using the method reported elsewhere 165 (Choi et al., 2012) with slight modifications. Frozen muscles were cut into 10 µm thick 166 transverse sections using a cryomicrotome (CM1860, Leica Biosystems Inc., USA) at -20°C. 167 Each sample was mounted on $76 \times 26 \times 1$ mm adhesive microscope slides (HistoBond[®], Paul 168 Marienfeld GmbH & Co. KG, Lauda-Königshofen, Germany), with a coverslip (22×22) mm, 169 and coated with a drop of aqueous mounting medium (S3023, Dako, Carpinteria, CA, USA) 170 with a 22×22 mm coverslip (100 Deckglaser, Menzel-Glaser). A fluorescent microscope 171 (BX51, Olympus, Tokyo, Japan) with a DP72 digital camera was used to display and 172 photograph all the samples (Olympus). Muscle fiber density (fiber number/4000 μ m²) were 173 determined using Photoshop CC (Adobe, California, USA). 174

175

176 Field emission scanning electron microscopy (FE-SEM)

FE-SEM was carried out at Cheonnam National University, Yeosu Campus to observe the sarcomere structure and muscle fiber size in the extrudates. The extrudates were split into $1 \times 1 \times 0.2$ cm dimensions. For analysis, each test extrudate was cut in both the transverse and longitudinal directions. Double-sided carbon tape was used to adhere the test samples to aluminum stubs. Meat cuts from different treatments were pre-fixed with 2.5% glutaraldehyde
for 4 h at 4°C. Subsequently, one percent osmium tetroxide was used to post-fix the samples
at 4°C for 2 h (dark condition). The samples were soaked thoroughly in PBS and washed in a
gradient of ethanol (30% to 99%), air-drying each wash. For 1 min, a platinum coating was
applied at 10 mA. Imaging was carried out using a Sigma 500 FE-SEM (Zeiss, Oberkochen,
Germany).

187

188 Statistical analysis

189 Statistical analysis was conducted using the GLM procedure (SAS, 2003). A significance 190 test between the results was performed using the Student-Newman-Keuls test as a multiple 191 assay technique (p<0.05). The analysis results are presented as the standard errors of the mean 192 and the least square means.

193

194 **Results and discussions**

195 pH, fat, and cholesterol content

Table 1 lists the pH, fat, and cholesterol content of A, B, W, and/broiler from the breast 196 and thigh meat at 5 and 12 wk respectively. At 5 wk, there were no significant variations in pH, 197 fat, or cholesterol content among the tested groups (W, A, and B) in the breast meat (p > 0.05). 198 On the other hand, at 12 wk, the pH and fat content were significantly higher in the broiler than 199 200 A and B, while the native chicken strains had a significantly higher cholesterol content than the broilers (p < 0.05). The lower pH in new native chicken strains (A and B) at 12 wk might be 201 due to the faster energy metabolism, which probably enhances glycogen storage in the muscle 202 203 (Fernandez et al., 2001). Typically, the composition of the body and muscle changes with the animal's age; the protein and fat content increase, and the moisture content decreases (Carmack 204 et al., 2020). Generally, the cholesterol content in slow-growing chickens is lower than other 205

206 meat-type chickens (Ali et al., 2019; Jaturasitha et al., 2008), which does not agree with the present study. The higher cholesterol levels in the new native chickens might be due to the 207 same diet without pasture and the ultimate deposition of additional saturated fatty acids (SFAs) 208 209 than broilers in the breast at 12 wk (Molee et al., 2012). The cholesterol level, however, varies according to the contents of body fat, skin fat, and bone marrow (Demos and Mandigo, 1995). 210 In thigh meat, at 5 wk, the new native chicken strains had a significantly higher pH than 211 W (p < 0.05), which might be due to lower lactate production with a lower buffering capacity 212 by the native chicken muscle type I in early stage of production that is dominant in thigh meat 213 (Listrat et al., 2016). On the other hand, the pH of broiler meat was significantly higher than 214 that of the new native chicken strains at 12 wk. This may be due to more aggressive behavior 215 by the new native chicken strains, which induces high levels of stress, which, in turn, draws 216 217 more glycogen into use. As a result, the postmortem glycolysis process is affected significantly, resulting in high lactic acid accumulation and a low pH in the meat (Jaturasitha et al., 2002). 218 Genetics - breed lines, gender, how animals are kept, transport, lairage conditions and time -219 pre-slaughter tension, method of slaughter, technical parameters and postmortem handling, and 220 meat storage time, all affect the pH of meat (Ristic and Damme, 2013). Furthermore, because 221 the pH of meat varies according to the muscle type of the same animal, it is important to 222 standardize the muscle area where the pH would be measured. This study analyzed the pH, fat, 223 224 and cholesterol content in the two new lines of native chicken strains (A and B) with slaughter 225 ages of 5 and 12 wk respectively (data are not shown). The pH of the breast meat was affected by age, whereas the fat and cholesterol contents were unaffected by the strain or age. Regarding 226 muscle pH, older birds had a lower pH, which concurs with Glamoclija et al. (Glamoclija et al., 227 228 2015). On the other hand, the pH and fat content in thigh meat were affected by age. Thus, native chicken strains provide valuable information on the meat quality traits seeking consumer 229 acceptance from the breast and thigh distinctively. 230

231 Fatty acid compositions

Table 2 lists the fatty acid compositions of A, B, W, and/broiler from the breast and thigh 232 meat at 5 and 12 wk respectively. At 5 wk, the breast meat of the new strains contained a higher 233 content of linoleic acid (C18:2), which is important for a similar flavor to cooked chicken meat 234 flavor (Rikimaru and Takahashi, 2010). White semi-broiler (W) is traditionally used for 235 making the unique cuisine, samgyetang, in Korea (Nam et al., 2010). The new native chicken 236 strains, however, outperformed the broilers in terms of eicosatrienoic acid (C20:3), ARA, EPA, 237 238 DHA, and nervonic acid at 12 wk (p < 0.05). This result is consistent with the findings reported by Rikimaru and Takahashi, (2010), who analyzed the fatty acid compositions of Hinai-jidori 239 chickens and broiler in Japan. Jayasena et al. (2013) reported that Korean native chickens 240 contained high levels of EPA and DHA compared to broilers whose contents resembles ω-3 241 PUFAs. On the other hand, the new native chicken strains had significantly higher SFAs and 242 PUFAs contents (p < 0.05) than the broilers (Jayasena et al., 2013) at 12 wk. Moreover, the new 243 native chicken strains had a lower ω -6/ ω -3 PUFA ratio than the broiler chickens at 12 wk, 244 which is desirable in foods of an animal origin (p < 0.05). Regardless of the muscles, at 12 wk, 245 246 however, the new native strains were enriched with the most favorable fatty acids. Omega-3 fatty acids compared to the broiler (p < 0.05) has a key function in the human body. The lower 247 ω -6/ ω -3 PUFA ratio of the new native chicken meat was due to the higher total ω -3 PUFAs 248 249 levels in the new native chicken strains and is associated with the breed or line effects with different genetic makeup (Ali et al., 2021; Enser et al., 2000). Compared to the broilers, the 250 PUFA/SFA (P/S) ratio has a favorable balance between the ω -6 and ω -3 PUFA, and new native 251 chicken strains had significantly higher levels than the broilers in the present study because of 252 the higher PUFAs content. Thus, the breast meat from young new native chicken strain could 253 254 be replaced with white semi-broiler for samgyetang and old new native chicken strains perform

better than broiler because of the ARA, DHA, nervonic acid, and omega-3 fatty acids with a lower ω -6/ ω -3 PUFA ratio.

At 5 wk, the thigh meat did not show any significant variations in fatty acid composition 257 among the W, A, and B tested groups (p>0.05. On the other hand, the older (12 wk) native 258 chicken strains had significantly higher levels of ARA, DHA, nervonic acid, and frequently 259 omega-3 fatty acids than the broilers (p < 0.05), which is consistent with Rikimaru and 260 Takahashi, (2010). New native chicken strains contained a higher total of SFAs than broiler 261 (p < 0.05). Nevertheless, compared to broilers, the ω -6/ ω -3 PUFA ratio showed a similar trend 262 to breast meat. These results suggest that the ARA content in new native chickens meat appears 263 to be a characteristic feature of meat from new native chicken strains developed recently in 264 Korea. The additional analysis assigned with two new native chickens showed that the age of 265 266 the birds had greater effects on the fatty acid profile than the line of the birds. On the other hand, the impact of both varied between the breast and thigh meat. The ratio of ω -6/ ω -3 PUFA 267 in the breast and thigh was lower in older chickens (p < 0.001), which is important for foods 268 and was due to an increase in the ω -3 PUFAs content (Popova et al., 2016). The fatty acid 269 composition could depend on age, line, and interactions within or between the breast and thigh 270 at different slaughter ages (Popova et al., 2016). 271

272

273 Antioxidant dipeptides (carnosine and anserine content)

Table 3 lists the antioxidant di-peptides in breast and thigh meat from two slaughter ages. Anserine is one of the predominating histidine di-peptides in new native chicken compared to broilers (Ali et al., 2019). The new native chicken strains had higher anserine levels than W or broilers at 5 and 12 wk, respectively (p<0.05), which is in line with Ali et al. (Ali et al., 2019), who compared the three new native chicken strains with broilers as well as commercial native chickens. In breast meat, at both slaughter ages, the carnosine content in W or broiler 280 was significantly higher than in the new native chicken strains, which is in agreement with Jung et al. (2013). The effect of the meat type on the anserine content was the same as the effect 281 of the meat type on the carnosine content. Because anserine acts as a physicochemical buffer 282 283 against the proton output by anaerobic glycolysis in different muscle traits, the new native chickens had higher anserine levels than the broilers (Dunnett and Harris, 1995). White muscle 284 tissue had a higher carnosine concentration than red muscle tissue, which tends to generate 285 energy-rich phosphate ester under anaerobic conditions (Boldyrev et al., 2004). The anserine 286 and carnosine content in thigh meat showed a similar fashion to breast meat. As a part of 287 additional analysis, the carnosine and anserine contents in the breast of birds were influenced 288 by their age rather than their lines. The carnosine levels were higher in the young birds, while 289 290 the anserine levels were higher in the older birds (p < 0.001). On the other hand, the carnosine 291 levels in the thigh meat were unaffected by age, but the anserine levels increased with age (p < 0.001). According to Chan et al. (1994), the muscle anserine content is affected by the breed, 292 muscle type, and animal age. 293

294

295 Muscle fiber density

Table 4 lists the muscle fiber density of the breast meat from W, and broilers with A and 296 B at 5 and 12 wk respectively. Regardless of the slaughter age, A and B had a higher fiber 297 298 density than the W or broiler. This study shows that a higher muscle fiber density in new native 299 chicken strains was in good agreement with a study reported by Koomkrong et al. (2015) conducted on Thai native chickens and broilers. Different muscle types, ages, breeds, and 300 muscle fiber sizes can influence the number of fibers in meat. Chicken and pigs have 301 302 experienced improvements in both the number and size of muscle fibers due to selection for an increased growth rate (Chen et al., 2007). The total amount, density, and composition are 303 essential histochemical attributes that influence the fresh or cooked meat during muscle to meat 304

305 conversion (Joo and Kim, 2011). Intrinsic and extrinsic factors can control muscle fiber biochemical and structural characteristics independently to increase the production efficiency 306 and enhance meat quality (Listrat et al., 2016). The morphological, contractile, and metabolic 307 308 properties of these muscle fibers define the characteristics of muscle fibers (Joo et al., 2013). Furthermore, the number of muscle fibers is determined genetically before birth, and only the 309 length and cross-sectional area of the muscle fibers increase with age (Wigmore and Stickland, 310 1983). The influences of the line and age in the two lines of the new native chickens at 5 and 311 12 wk respectively and were examined further. The findings show that age has a larger effect 312 on the muscle fiber profile than the line (p < 0.001). The muscle fiber density of the older native 313 chickens was lower than that of the young birds (p < 0.001). Furthermore, regarding the muscle 314 fiber region, the genotype effect interacts with age (Chen et al., 2007). Therefore, breed and 315 age can have distinct and different effects on carcass performance and meat quality in 316 commercial broilers or white semi-broilers and indigenous chickens. 317

318

319 Field emission electron microscopy (FE-SEM)

Figs. 1 and 2 present FE-SEM images of the size and compactness (cross-section) of the 320 muscle structures with the degree of the sarcomere pattern (longitudinal) of the breast meat 321 excised from W and/broilers, A, and B at 5 and 12 wk respectively. Regardless of the slaughter 322 age, the muscle fiber size, which is positively correlated with the carcass weight, was larger in 323 324 W or broilers than the new native chicken strains; A and B. A previous study showed that the imported breed chickens have a larger fiber size in the breast and thigh muscles than Thai native 325 chickens (Jaturasitha et al., 2008). The larger body weight of chickens is based on the larger 326 327 muscle fiber diameter and area and lower muscle fiber density (Chen et al., 2007). The new native chicken strains arrange the muscle fiber in a smaller pattern and more compactness, 328 making the muscle tough. A denser muscle fiber was noted in the 5-wk-old birds. These 329

330 structural differences at different lines and ages might be due to the differences in line, age, rate of rigor onset, and degree of sarcomere shortening (Smith and Fletcher, 1988). The cross-331 section of a muscle fiber increases more than that of the endomysium and perimysium 332 connective tissue during breast meat production in modern broiler breeds (Dransfield and 333 Sosnicki, 1999). This suggests that the muscles selected for rapid growth have outgrown their 334 life support systems, resulting in muscle injury (Swatland, 1990). Furthermore, with a fixed 335 336 focus (1 μ m), more sarcomere patterns were noted in new native chicken strains than W or broilers for both ages and clearly defined as older birds. On the other hand, the meat 337 tenderization in native chicken strains was attributed to a shorter sarcomere length (Van Laack 338 et al., 2000). 339

340

341 Conclusion

342 The new native chicken strains (A and B) had distinct quality features compared to white semi-broilers or broilers. The breast and thigh meat showed distinct meat quality traits. On the 343 other hand, the meat from the new native chicken strains had some unique meat quality traits 344 345 and exhibited more advantages over commercial birds (broilers or white semi-broilers), due primarily to the ARA content, which is related to the palatability of meat. In addition, the new 346 native chicken strains also performed better in terms of omega-3 fatty acids, DHA, nervonic 347 acid, ω -6/ ω -3 PUFA ratio, and most abundant antioxidant dipeptide, anserine. Thus, the current 348 investigation provides valuable information for chicken meat from new native strains, which 349 will influence the consumption of different chicken meat with desirable characteristics. Further 350 research will be needed to determine the relationship between those compounds and the 351 palatability of new native chicken strains. 352

353

354 **Conflict of Interest**

355 The authors declare no potential conflict of interest.

356

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361

362 Author Contributions

- 363 Conceptualization: Nam KC. Data curation: Nam KC, Ali M. Formal analysis: Park JY,
- Lee SY. Methodology: Park JY, Lee SY. Validation: Nam KC. Writing original draft: Ali M.
- 365 Writing review & editing: Ali M, Lee SY, Park JY, Nam KC.

366

Ethics Approval

- 368 This article does not require IRB/IACUC approval because there are no human and
- 369 animal participants.

Tables and figures

T.	5 wk					12 wk		CEN (1)
Items	W*	А	В	SEM ¹⁾	Broiler*	А	В	-SEM ¹⁾
Breast								
pН	5.85	5.83	5.88	0.02	6.24 ^a	5.75 ^b	5.73 ^b	0.02
Fat (%)	1.13	1.14	1.15	0.13	2.34 ^a	1.18 ^b	1.29 ^b	0.16
Cholesterol (mg/100g)	82.82	87.23	84.1	6.35	42.31 ^b	73.68 ^a	77.39 ^a	4.83
Thigh								
pН	6.40 ^b	6.54 ^a	6.55 ^a	0.02	6.62 ^a	6.34 ^b	6.30 ^b	0.02
Fat (%)	3.67	4.16	3.35	0.23	3.61	3.13	3.02	0.31
Cholesterol (mg/100g)	69.71	58.80	61.84	5.89	56.09	71.38	62.93	4.16
^{a-b} Values with				within the	same row di	ffer signif	icantly (p	< 0.05)
¹⁾ SEM: standar								
Star (*) indicat	tes a simila	r weight t	o native c	chickens.				
					~			

Table 1. pH, fat, and cholesterol of the breast and thigh meat in white semi-broiler (W), and/broiler with two new native chicken strains (A and B) according to slaughter age

Fatty acid	5 wk			– SEM ¹⁾		12 wk			
Fatty actu	W*	А	В		Broiler*	А	В	SEM ¹⁾	
Breast									
C12.0	n.d.	n.d.	n.d.		0.06 ^a	0.00 ^b	0.00 ^b	0.00	
C14:0	0.42	0.39	0.43	0.03	0.76^{a}	0.50^{b}	0.40 ^c	0.03	
C16:1	2.5	2.69	2.43	0.19	4.71 ^a	2.54 ^b	2.14 ^b	0.21	
C18:0	10.52	10.12	10.16	0.23	6.87 ^b	9.30 ^a	9.84 ^a	0.23	
C18:1	27.97	28.42	27.81	0.80	38.35 ^a	28.10 ^b	26.52 ^b	0.87	
C18:2	17.40 ^b	17.67 ^{ab}	17.92 ^a	0.12	19.19 ^a	15.49 ^b	15.28 ^b	0.31	
C18:3	0.36 ^a	0.29 ^b	0.30 ^b	0.01	0.52 ^a	0.23 ^b	0.22 ^b	0.02	
C20:2	0.59 ^a	0.43 ^b	0.43 ^b	0.03	0.41 ^a	0.30 ^c	0.36 ^b	0.01	
C20:3	1.55	1.38	1.4	0.07	0.68 ^c	1.00 ^b	1.19 ^a	0.05	
C20:4	7.67	7.17	7.18	0.36	2.32 ^b	8.81 ^a	9.70 ^a	0.55	
C20:5	0.22	0.24	0.23	0.02	0.13 ^b	0.17 ^{ab}	0.18 ^a	0.01	
C22:6	0.86	0.83	0.88	0.05	0.14^{b}	1.00 ^a	1.07 ^a	0.07	
C24:1	1.67	1.58	1.59	0.08	0.62 ^c	2.05 ^b	2.38 ^a	0.11	
$\sum SFA^{2)}$	32.40	32.32	32.5	0.14	29.74 ^b	32.16 ^a	32.37 ^a	0.26	
$\sum UFA^{3)}$	60.79	60.69	60.17	0.45	67.05 ^a	59.68 ^b	59.03 ^b	0.54	
$\sum MUFA^{4)}$	32.14	32.69	31.83	0.91	43.67 ^a	32.69 ^b	31.04 ^b	0.96	
$\sum PUFA^{5}$	28.65	27.99	28.35	0.53	23.38 ^b	26.99 ^a	27.99 ^a	0.51	
UFA/SFA	1.88	1.88	1.85	0.01	2.26^{a}	1.86 ^b	1.82 ^b	0.03	
P/S	0.88	0.87	0.87	0.01	0.79 ^b	0.84 ^a	0.86^{a}	0.016	
∑ω-6	25.66	25.26	25.53	0.43	21.91 ^a	24.60 ^a	25.33 ^b	0.44	
$\sum \omega - 3$	2.99	2.73	2.81	0.12	1.47 ^b	2.40 ^a	2.66 ^a	0.10	
ω-6/ω-3	8.59	9.25	9.27	0.33	15.01 ^a	10.36 ^b	9.61 ^b	0.39	
Thigh									
C12:0	n.d.	n.d.	n.d.		0.06 ^a	0.05 ^b	0.04 ^c	0.00	
C18:0	7.79	7.26	7.58	0.16	6.73 ^b	7.99 ^a	7.53 ^a	0.18	
C18:1	37.23	38.15	37.10	0.54	38.55 ^a	35.05 ^b	35.81 ^b	0.41	
C18:2	18.28	17.97	18.08	0.2	19.74 ^a	18.02 ^b	18.00 ^b	0.25	
C18:3	0.34	0.34	0.34	0.01	0.49 ^a	0.39 ^b	0.41 ^b	0.01	
C20:2	0.20 ^a	0.17 ^b	0.20 ^a	0.01	0.28 ^a	0.22 ^b	0.23 ^b	0.01	
C20:3	0.49	0.40	0.46	0.02	0.50	0.43	0.45	0.02	
C20:4	3.15	2.61	3.05	0.19	2.03 ^b	4.49 ^a	4.00 ^a	0.17	
C20:5	0.06	0.05	0.04	0.00	0.09 ^a	0.06^{b}	0.06 ^b	0.00	
C22:6	0.26	0.23	0.3	0.02	0.11 ^c	0.44^{a}	0.34 ^b	0.01	
022.0									

Table 2. Fatty acid compositions (%) of the breast and thigh meat in white semi-broiler (W),
and/broiler with two new native chicken strains (A and B) according to slaughter age

$\sum SFA^{2)}$	30.93	30.88	30.85	0.16	29.65 ^b	31.01 ^a	30.75 ^a	0.19
$\sum UFA^{3)}$	65.79	65.84	65.26	0.26	67.64 ^a	65.09 ^c	65.68 ^b	0.20
$\sum MUFA^{4)}$	43.01	44.08	42.79	0.56	44.42 ^a	41.05 ^b	42.20 ^b	0.49
$\sum PUFA^{5)}$	22.78	21.77	22.47	0.39	23.22	24.04	23.48	0.42
UFA/SFA	2.13	2.13	2.12	0.01	2.28^{a}	2.10 ^b	2.14 ^b	0.02
P/S	0.74	0.71	0.73	0.014	0.78	0.78	0.76	0.015
$\sum \omega - 6^{6}$	21.63	20.74	21.33	0.36	22.04	22.73	22.22	0.40
$\sum \omega - 3^{7}$	1.15	1.03	1.14	0.04	1.18 ^b	1.32 ^a	1.26 ^{ab}	0.03
ω-6/ω-3	18.87	20.26	18.80	0.51	18.75 ^a	17.30 ^b	17.64 ^b	0.34

- ^{a-c} Values with different superscripts letters within the same row differ significantly (p < 0.05). Star (*) indicates a similar weight to native chickens.
- ¹⁾ SEM: standard error of the means (n=18).
- ²⁾ SFA: saturated fatty acid.
- ³⁾ UFA: unsaturated fatty acid.
- ⁴⁾ MUFA: monounsaturated fatty acid.
- ⁵⁾ PUFA: polyunsaturated fatty acid.
- ⁶⁾ ω -6: Sum of linoleic acid (18:2), arachidonic acid (20:4), and eicosadienoic acid (20:2).
- ⁷⁾ ω-3: Sum of linolenic acid (18:3), EPA (20:5), DHA (22:6), and eicosatrienoic acid (20:3).

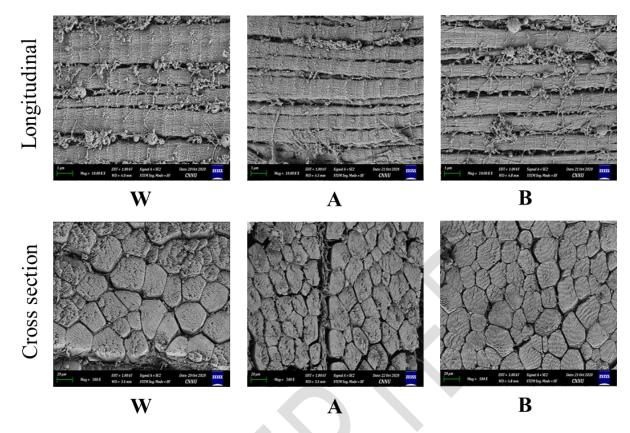
Table 3. Antioxidant di-peptides (mg/100 g) of the breast and thigh meat in white semi-broiler

417 (W), and/broiler with two new native chicken strains (A and B) according to slaughter age

Antioxidant		5 wk		SEM ¹⁾		12 wk		SEM
di-peptides	W*	А	В	SEM	Broiler*	А	В	SEM
Breast								
Carnosine	281.79 ^a	248.19 ^b	242.47 ^b	8.84	260.01 ^a	177.23 ^b	210.46 ^b	11.8
Anserine	544.64 ^b	638.31 ^a	607.79 ^a	10.35	335.08 ^b	693.89 ^a	681.30 ^a	15.4
Thigh								
Carnosine	148.95 ^a	111.89 ^b	109.52 ^b	4.80	176.38 ^a		119.20 ^b	4.3
Anserine	229.36 ^b		259.45 ^a	4.87	185.60 ^b		316.04 ^a	9.8
^{a-c} Values with ¹⁾ SEM: standar				ithin the	same row d	lifter sign	ificantly (<i>p</i> <0.0
Star (*) indicat				hickens.				
()		8						

Table 4. Muscle fiber density of the breast meat in white semi-broiler (W), and/broiler, with
two new native chicken strains (A and B) according to slaughter age

Item		5 wk SEM ¹) 12 wk			SEM ¹⁾			
Item	W*	А	В		Broiler*	А	В	
Muscle fiber density ²⁾	47.0 ^b	66.0 ^a	66.4 ^a	0.14	21.2 ^c	39.4 ^a	30.8 ^b	0.67
^{a-c} Values with d				within the sa	ame row dif	fer signi	ficantly	(<i>p</i> <0.05).
¹⁾ SEM: standard				m^2) of h_{max}	at most			
²⁾ Muscle fiber de Star (*) indicate					ist meat.			
2002 ()								



461 Fig. 1. Field emission scanning electron micrographs of longitudinal and cross-section of breast
462 meat from W, A, and B at 5 wk

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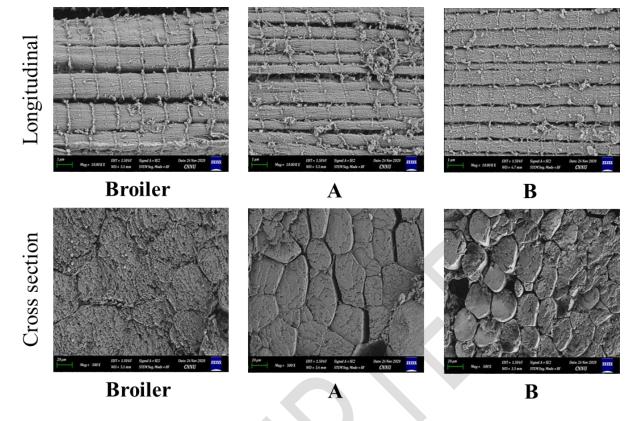
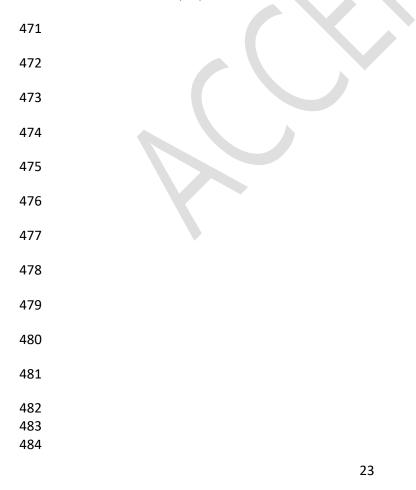


Fig. 2. Field emission scanning electron micrographs of longitudinal and cross-section of breast
 meat from broiler, A, and B at 12 wk



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