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The Quality Characteristics of Ready-to-Eat Empal Gentong affected by meat pre-cooking

Abstract

The purpose of this research was to examine the effectiveness of pre-cooking treatments on the quality characteristics of ready-to-eat (RTE) empal gentong. Raw beef meat was precooked in water bath at 90°C for 0 min (C), 10 min (T1), 20 min (T2), and 30 min (T3) prior to retorting process at 121°C and pressure at 0,7 bars. Results showed that pre-cooking treatments in all treated samples could reduce fat contents in empal gentong's meat by 0.02% (T1), 0.28% (T2), and 1.13% (T3) respectively. Highest precooking time tends to increase the pH and CIE a* values. However, CIE b* values, water holding capacity (WHC), and sensory analysis were not affected by pre-cooking duration which must have been affected by sterilization process after pre-cooking. In conclusion, pre-cooking treatment before sterilization in producing empal gentong is a probable technique to reduce its fat content and improve its physical quality. A specific treatment at 90°C for 10 min is recommended to achieve optimum quality of ready-to-eat empal gentong's meat.

Keywords: pre-cooking, meat, ready-to-eat, empal gentong, quality characteristic

Introduction

Interest in traditional food products has grown in both developed and developing countries (Anders and Caswell 2009). As a country with diverse cultures and traditions, Indonesia has a variety of traditional foods (Rianti et al. 2018). Empal gentong, a traditional food originating in Cirebon, Indonesia, is meat prepared with mixed spices and coconut milk. However, consumers nowadays complain about fat droplets in the broth and the short shelf life of the product. Therefore, retort packaging, which involves sterilization at high temperatures, is used to produce ready-to-eat (RTE) empal gentong.

35 The quality of RTE empal gentong meat was the focus of this study. Some small and
36 medium level industries still use the conventional method of pre-cooking without a
37 standard. However, several recent studies have aimed to reduce the fat content and
38 improve the visual appeal of meat products. Triyannanto and Lee (2015) showed that pre-
39 cooking successfully improves the quality of Korean ginseng chicken soup, as judged by
40 consumer acceptance. Furthermore, Manheem et al. (2013) reported that a cheap and
41 simple pre-cooking process is important for extending the shelf life of food products.
42 Accordingly, our study aimed to identify the optimal pre-cooking method for RTE empal
43 gentong by evaluating the quality characteristics of the meat prepared using various pre-
44 cooking treatments.

46 Materials and Methods

47 Meat preparation and precooking treatment

48 Fresh beef meat was purchased from the local butcher's market in Yogyakarta City of
49 Indonesia, and immediately brought to the laboratory. The fresh beef meat (*longissimus*
50 *dorsi*) was cut into cubes with a size of 3x3x3 cm (LxWxH) to be prepared for pre-
51 cooking treatment. The meat samples were then packed with sealable PE plastics bag.
52 The precooking process was carried out by heating the meat samples in the water bath at
53 a temperature of 90°C. There were four group treatment of pre-cooking time namely
54 control/without pre-cooking (C), 10 min (T1), 20 min (T2) and 30 min (T3) of pre-
55 cooking time with five replications. The curry was separately prepared by mixing the
56 coconut milk, spices, and hot water. The curry was heated at a temperature of 80-90°C
57 for 45 min.

59 RTE Empal gentong preparation

60 A total of 50 g meat cubes was introduced to multilayer retort pouch with a specific
61 layer arrangement of PET / ALU / ONU / CPP, 16.0 cm x 22.9 cm (WxH) size. About
62 300 ml of hot curry was poured into the pouch, then was sealed by using a continuous
63 sealer machine. Afterward, the sterilization process was carried out using a retort machine
64 which was operated by holding a pressure of 10.15 psi, 9 min until sterility value is
65 obtained. After sterilization, a cooling process was carried out in room temperature water
66 at 22-25°C for 10 min. The RTE empal gentong samples were then analyzed.

67

68 pH value

69 Ten grams of empal gentong's meat was chopped and then transferred into 40 mL of
70 distilled water, homogenized at 10.000 rpm for 60 s using a homogenizer. The pH values
71 were measured using a pH meter attached with an electrode (Orion Star A111 Benchtop,
72 Thermo Fisher Scientific Inc, Singapore). The pH value was performed in triplicate per
73 treatment (Muhlisin et al. 2013).

74

75 Tenderness

76 Samples of empal gentong's meat with a thickness of 0.5 cm and 1.5 cm width were
77 placed on the Warner-Bratzel instrument (Soeparno 2015)

78

79 Water holding capacity

80 The analysis of the water holding capacity (WHC) in this research using the method of
81 (Hamm 1972). Samples in amount of 0.3 gr placed on filter paper and pressed between 2
82 glass plates, and then given 35 kg load for 5 min. The area which absorbed water was
83 then counted with planimeter. WHC then calculated with the following formula:

84

$$\text{mgH}_2\text{O} = \frac{\text{wet area (cm}^2\text{)} - 8}{0,0948}$$

85
$$\% \text{ free water} = \frac{mgH_2O}{weightsample(mg)} \times 100\%$$

86 The sample used for water content assay was 1 g. Weighed samples then inserted into
87 filter paper and oven dried at 105°C for 24 h (Soeparno 2015).

88
$$WHC = \frac{x + y - z}{x} \times 100\%$$

89
$$\%WHC = TWC - \% \text{ free water}$$

90 Where X = Sample weight; Y = filter paper weight; Z = Sample weight + filter paper
91 weight after being oven; and TWC = Total Water Content.

92

93 Cooking loss

94 The analysis of the cooking loss in this research using the method of Bouton et al.,
95 (1972). The meat was cut in the direction of the fiber and weighed as much as 25 g.
96 Afterwards, the meat was put in polyethylene plastic and packed with a vacuum machine.
97 The meat was cooked in a water bath at 90°C for 0 min (C), 10 min (T1), 20 min (T2),
98 and 30 min (T3) min. The meat was then cooled and removed from the polyethylene
99 plastic and then wiped with a tissue and the final weight is weighed.

100
$$\text{Cooking Loss} = \frac{x - y}{x} \times 100\%$$

101 Where x = initial weight; y = final weight

102

103 Proximate analysis

104 Chemical analysis method for this research were water, fat, protein, and collagen
105 content by using a food scanner (FoodScan™ Meat Analyser; FOSS, Padova, Italy) with
106 NIRS (Near Infrared Reflectance Spectroscopy) technology. Thirty grams of sample were
107 grinded and checked in food scanner with a special petri dish. Samples checked in
108 triplication (Triyannanto et al. 2019)

109 Sensory analysis

110 Sensory analysis following the method described by Triyannanto and Lee, (2015). The
111 total of 11 male and female semi-trained panellists aged 17- 21 years conducted a sensory
112 analysis for RTE empal gentong. Sensory procedures were explained in detail to the
113 panellists before conducting a sensory test. A pack questionnaire was given to be filled
114 during a sensory analysis. Every sample was labelled with 3 different numbers to decline
115 the subjective score possibility. To support the sensory analysis lamp room with a 1,200-
116 lux brightness were applied. Panellists are required to rinse their mouth after the analysis
117 for each different sample. These procedures were designed to avoid cross-contamination
118 of the sensory characteristics in each sample. Furthermore, the panellist was obliged to
119 fill the questionnaire that has been provided. Sensory analysis in this research was
120 contained of four parameters namely, color, tenderness, taste, texture, and flavour.
121 Parameter scales were set at; 5: very like, 4: Like, 3: plain 2: dislike, and 1: very dislikes.

123 Statistical analysis

124 SPSS Statistics (version 25.0; (IBM, 2017) for Windows Evaluation Version was used
125 to analyze all data. The data were analyzed using one way analysis of variance and
126 Duncan's multiple range test for significant differences ($p < 0.05$).

128 Results and Discussion

129 pH value

130 Table 1 shows that pre-cooking time significantly affected the pH value of the meat (p
131 < 0.05). The pH value was 6.31 in the control and tended to increase with longer pre-
132 cooking times. The pH values in this study might have been affected by the heating
133 process, which causes amino acids to lose their carboxyl groups. A decrease in the number
134 of acidic groups was also observed by Hamm and Deatherage (1960), who showed that

135 ground longissimus dorsi muscle lost almost one-third of its carboxyl groups when heated
136 at 20-70°C for 30 min in a water bath contained by a covered metal vessel.

137

138 Water-holding capacity

139 Table 1 shows that the pre-cooking time did not significantly affect the water-holding
140 capacity of the meat ($p > 0.05$). The water-holding capacity was dependent on the amount
141 of denaturation of the meat protein. The absence of a significant effect was possibly
142 caused by the complete denaturation of protein during the sterilization process at 121°C,
143 which resulted in a constant water-holding capacity among all treatments. High pressure
144 thermal processing after pre-cooking results in complete protein denaturation. In
145 accordance with this, Sun et al. (2016) reported that in beef, pork, and chicken,
146 commercial sterilization at 121°C for 10 min leads to protein-bound water but does not
147 significantly affect the protein and fat content in beef and pork. Moreover, Soeparno
148 (2015) showed that myofibril protein coagulates at 30°C and completely denatures at
149 55°C, which is lower than the commercial sterilization temperature. Furthermore, Gómez
150 et al. (2020) reported that high pressure and temperature do not significantly affect the
151 cooking loss rate or water-holding capacity.

152

153 Tenderness

154 The tenderness of the meat was measured by determining the content of connective
155 tissue, such as collagen. As shown in Table 2, decreased penetrometer values indicated
156 that tenderness increased significantly ($p < 0.05$). Pre-cooking produced penetrometer
157 values of 4.26 kg/cm² (T1), 4.23 kg/cm² (T2), and 4.13 kg/cm² (T3), which were lower
158 than those of the control (6.40 kg/cm²). A lower penetrometer value objectively shows
159 that less energy and pressure are required for chewing. Collagen hydrolysis during pre-

160 cooking resulted in increased tenderness. Lawrie and Ledward (2006) reported that
161 cooking affects meat structure, softening the connective tissue by converting collagen
162 into gelatin. Moreover, Soeparno (2015) stated that tenderness reflects the amount of
163 collagen present and that long boiling times cause changes in the structure of muscle
164 proteins, especially actin and myosin. The breakdown of actin and myosin can influence
165 the mechanical strength of connective tissue (Bouton and Harris, 1972).

166

167 Cooking loss

168 The cooking loss observed in each pre-cooking condition is presented in Table 2. An
169 extrinsic factor that affected cooking loss was pre-cooking duration. Meat subjected to
170 longer pre-cooking treatments tended to exhibit significantly greater cooking losses than
171 those subjected to shorter treatments. Meat in the T3 group, pre-cooked for 30 min,
172 exhibited the greatest cooking loss. This loss might consist of water and other water-
173 soluble components, such as proteins. High pre-cooking temperatures up to 90°C
174 decreased the initial weight of the empal gentong meat by almost half. This result was in
175 accordance with that found by Tornberg (2005), who stated that the greatest cooking loss
176 in beef occurs at 60-80°C, which is lower than the pre-cooking temperature used in our
177 study. Hearne et al. (1978) also reported that higher endpoint temperatures result in
178 greater cooking loss in bovine semitendinosus meat.

179

180 Instrumental color

181 The instrumental color values, CIE L* (lightness), a* (redness), and b* (yellowness),
182 are presented in Table 1. The CIE L* and a* values of RTE empal gentong meat in the
183 T2 group were lower than those in the T1 group, indicating that these values tended to
184 decline with longer pre-cooking times ($p > 0.05$). However, the highest values were

185 observed in meat pre-cooked for 30 min (T3) ($p < 0.05$). As reported by Muhlisin et al.
186 (2013), chuncheon dalkalbi meat with a lower CIE L* value exhibits a darker color. The
187 effect of pre-cooking time on the CIE L* and a* values of empal gentong meat in this
188 study was not clear. Certain ingredients of RTE empal gentong, such as turmeric, ginger,
189 and other herbs, which naturally tend to be yellow in color, might have been responsible
190 for the CIE L* and a* values during processing. Longer pre-cooking time had no effect
191 on the CIE a* value ($p > 0.05$). It seemed that sterilization at 121 °C was responsible for
192 more defects than the pre-cooking duration, which produced a non-significant CIE b*
193 value. Myoglobin, responsible for the red color of meat, turns grayish brown at 75°C
194 (Hunt et al., 1999), which is lower than the pre-cooking temperature of 90°C and far lower
195 than the sterilization temperature of 121°C used in this study. Moriyama and Takeda
196 (2010) reported that myoglobin is mostly destroyed at 70-100°C.

197

198 Proximate composition

199 Table 2 shows the proximate composition of RTE empal gentong subjected to the
200 various pre-cooking times. Significant differences in moisture, protein, fat, and collagen
201 were observed between the control and experimental groups ($p < 0.05$). As shown in
202 Table 2, the moisture content of the control was 67.16 % (w/w), but that of the T1, T2,
203 and T3 groups was reduced by 1.54-5.92%. The lower moisture content of the meat
204 samples subjected to pre-cooking treatments might be related to the heat-induced
205 denaturation of myofibrillar protein, which can adversely affect the water-holding
206 capacity (Triyannanto and Lee 2015). This result is in accordance with the results shown
207 in Table 1, which indicated greater cooking loss with longer pre-cooking duration.

208 The crude protein content of the control was 23.93%, while that of the T1, T2, and T3
209 groups was reduced by 0.18-0.84%. Cooking at a high temperature for a long time causes

210 the protein content to decrease. Tornberg (2005) reported that the heating process results
211 in denaturation of myofibril proteins and changes in protein structure. In addition, the
212 soluble protein content decreases by approximately 90% as the meat temperature
213 increases from 23°C to 80°C (Murphy and Berrang, 2002). In accordance with this, the
214 decrease in protein content observed in our study paralleled the decrease in collagen
215 composition.

216 The crude fat content of the empal gentong control samples was 6.26%. Pre-cooking
217 significantly reduced this value by 0.02% (T1), 0.28% (T2), and 1.13% (T3). In this study,
218 pre-cooking prior to sterilization was a suitable way to reduce the fat content ($p < 0.05$).
219 However, the longer duration of pre-cooking required to achieve the decrease in fat
220 content might also decrease meat quality, manifested as a higher percentage of cooking
221 loss. Therefore, pre-cooking for 10 min could be a solution for producing RTE empal
222 gentong with a lower fat content as well as a higher proximate content. The RTE empal
223 gentong in the T1 group had better quality than that in the other pre-cooking treatment
224 groups, although there were no significant differences between the T1 group and the
225 control ($p > 0.05$). This result is in accordance with a prior study conducted by
226 Triyannanto and Lee (2015), who showed that precooking at 90°C for 10 min is an
227 effective way to improve the quality of Korean ginseng chicken soup.

228 Collagen content did not differ significantly between the control (2.53%) and T1 groups
229 (2.43%), while that of the T2 and T3 groups was significantly reduced by 0.10-0.47%.
230 There was no significant difference in collagen content between the T2 and T3 groups (p
231 > 0.05). The results showed that denaturation induced by pre-cooking tended to reduce
232 collagen levels. Tornberg (2005) reported that collagen denaturation occurred at 53-63°C
233 and that gelatin was formed with further heating. Some of the gelatin observed in this
234 research might dissolve in the empal gentong broth during the sterilization process.

235 As shown in Table 3, sensorial values in all treatment groups were not affected by pre-
236 cooking conditions ($p > 0.05$). The sterilization process, with a temperature of 121 °C and
237 pressure of 10.15 psi, probably had a greater effect than pre-cooking on sensorial values.
238 From this study, it could be concluded that sterilization at a high temperature and pressure
239 has a greater influence on all sensory qualities of empal gentong meat than the duration
240 of pre-cooking. As reported by Triyannanto and Lee (2015), heat exposure during
241 sterilization at 120 °C for 65 min had a greater impact than pre-cooking on ginseng
242 chicken soup products. However, although sterilization has a significant effect on sensory
243 qualities, it is necessary for producing RTE empal gentong that is free of spoilage-
244 inducing microbes and pathogens.

245

246 Conclusion

247 The current conventional empal gentong production without pre-cooking produce high
248 fat content, which tend to be unpleasant to consumer's perspective. Pre-cooking treatment
249 can be used in manufacturing RTE empal gentong to optimize its quality. Specific pre-
250 cooking condition at 90 °C for 10 min is recommended to maintain its proximate with a
251 lower fat content. This finding should be useful in the commercial production of RTE
252 empal gentong or other relevant products, giving an optional outcome with low fat but
253 high proximate content product as well as economically visible.

254

255 Conflicts of interest

256 The authors declare no potential conflict of interest

257

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262

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334 Tables

335

336 Table 1. pH value, Tenderness, WHC, Cooking Loss and Instrumental Color of Meat RTE

337 Empal Gentong Depending on Pre-cooking Conditions

Physical Parameters	Pre-cooking conditions			
	C (Not pre-cooked)	T1 (90°C/10 min)	T2 (90°C/20 min)	T3 (90°C/30 min)
pH value	6.31±0.01 ^b	6.33±0.06 ^b	6.34±0.01 ^b	6.41±0.02 ^a
Tenderness (kg/cm ²)	6.40±0.20 ^a	4.26±0.25 ^b	4.23±0.25 ^b	4.13±0.15 ^b
WHC ^{NS}	43.00±3.60	42.33±3.21	36.33±0.57	34.00±7.00
Cooking loss	-	39.00±1.73 ^b	41.67±1.15 ^b	46.67±2.30 ^a
CIE L*	15.53±0.25 ^b	14.30±0.95 ^c	13.66±0.15 ^c	20.20±0.00 ^a
a*	4.66±0.05 ^{ab}	4.43±0.25 ^b	3.90±0.88 ^b	5.43±0.11 ^a
b* ^{NS}	14.53±0.05	11.03±2.37	12.40±2.07	14.86±0.26

338 Results are expressed as mean±SD.

339 ^{a,b} Values within each row with different superscripts are significantly different (p<0.05).

340 ^{NS} Not significantly different (p>0.05)

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346

347 Table 2. Proximate Composition of RTE Empal Gentong's Meat Depending on Pre-
 348 cooking Conditions

Proximate Composition (%)	Pre-cooking conditions			
	C (Not pre- cooked)	T1 (90°C/10 min)	T2 (90°C/20 min)	T3 (90°C/30 min)
Moisture	67.16±0.10 ^a	65.62±0.38 ^b	65.49±0.04 ^b	61.24±0.21 ^c
Protein	23.98±0.14 ^a	23.80±0.27 ^a	23.57±0.50 ^{ab}	23.14±0.09 ^b
Fat	6.26±0.10 ^a	6.24±0.06 ^a	5.98±0.03 ^b	5.13±0.10 ^c
Collagen	2.53±0.29 ^a	2.43±0.08 ^a	2.25±0.17 ^b	2.06±0.06 ^b

349 Results are expressed as mean±SD.

350 ^{a,b,c}Values within each row with different superscripts are significantly different (p<0.05).

351 ^{NS} Not significantly different (p>0.05)

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364 Table 3. Sensory Characteristics of Meat RTE Empal Gentong Depending on Pre-cooking
 365 Conditions

Sensory analysis	Pre-cooking conditions			
	C (Not pre-cooked)	T1 (90°C/10 min)	T2 (90°C/20 min)	T3 (90°C/30 min)
Color ^{NS}	3.92±0.49	4.00±0.40	3.96±0.53	3.96±0.53
Texture ^{NS}	3.48±0.82	3.64±0.86	3.76±0.96	3.96±0.78
Flavor ^{NS}	3.72±0.61	3.72±0.61	3.76±0.59	3.80±0.76
Taste ^{NS}	3.40±0.86	3.44±0.86	3.48±0.87	3.40±0.76
Acceptability	3.60±0.64	3.64±0.70	3.72±0.54	3.64±0.63
NS				

366 Results are expressed as mean±SD.

367 ^{NS} Not significantly different (p>0.05)

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