ABSTRACT

Objective: The objective of this study is to investigate the effect of light-roasted green coffee bean extract (GCE) administration for 7 weeks on the improvement of metabolic profile, adiponectin level, homeostatic model assessment insulin resistance (HOMA-IR) index in metabolic syndrome (MS) rat model.

Methods: Adult male Sprague-Dawley rats were induced by a combination of high sucrose and high-fat diet for 8 weeks and streptozotocin injection in the 2nd week. The MS was confirmed by NCEP-ATP III criteria. They were divided into six weight-matched groups (n=5): normal control, MS, metformin and simvastatin-treated group (DMS), 100 and 200/body weight (bw) GCE (GCE 100 and GCE 200, respectively). The extracts were given through oral gavage daily for 7 weeks. The effect of GCE on body weight, serum glucose, triglyceride, (TG) and high-density lipoprotein (HDL) level was analyzed by colorimetric method. HOMA-IR index and adiponectin were analyzed by enzyme-linked immunosorbent assay methods.

Result: Fasting blood glucose, TG, and systolic blood pressure decreased significantly (p<0.05) in both GCE groups. Moreover, after 7 weeks, those parameters were significantly lower (p<0.05) compared to that of MS group. Only GCE 100 group that showed a significant decrease in HDL level. GCE 100 mg/bw and 200 mg/bw group showed significantly higher adiponectin level compared to that of MS and DMS group. Furthermore, GCE 100, GCE 200, and DMS group showed a significant lower HOMA-IR index compared to that of MS group.

Conclusion: 7 weeks GCE administration could decrease fasting blood glucose, profile lipid, blood pressure, and improved adiponectin level and HOMA-IR index.

Keywords: Metabolic syndrome, Green coffee bean extract, Adiponectin, Homeostatic model assessment insulin resistance.
index, lipid profile, fasting blood glucose level, and SBP in MS rat model.

METHODS
Extraction of green coffee bean
Coffee was extracted from light roasted robusta green coffee beans (Temanggung, Indonesia). Green coffee bean was obtained to sort out high-quality seed. Green coffee bean weighed 500 g was dried using a dryer cabinet on the temperature of 50°C for 8 h to obtain simplicia with 9-10% water content. The simplicia was mashed with a blender and then macerated by methanol to produce crude extract. The crude extract was filtered using a filter cloth to separate the liquid phase from the solid phase. The liquid phase was concentrated using rotary evaporator on temperature ±40°C. The concentrated liquid phase was partitioned using butanol, water, and acetylacetone. Finally, column chromatography was completed using silica gel as static phase, and the filtered product was evaporated.

Animal care and experimental protocol
25 Sprague-Dawley rats were purchased from the National Agency of Drug and Food Control, Indonesia. They were housed in standard cages and placed in a room where the temperature was maintained at 25±1°C, relative humidity at 50±1%, and the light at a 12 hrs light/dark cycle. During a 1-week acclimatization period, all rats consumed a normal pelit diet and tap water ad libitum. Afterward, they received high sucrose, HFD, and high sodium diet for 8 weeks and intraperitoneal streptozotocin injection (30 mg/kg body weight [bw]) in the 2nd week. The injection was repeated in the 3rd week with the same dose. The rats with blood glucose (>126 mg/dL), TG (>150 mg/dL), high SBP (≥140 mmHg), and reduced HDL levels (<40 mg/dL) was confirmed as MS rat based on NCEP-ATP III criteria [2]. Furthermore, they were divided into six weight-matched groups (n=5): The normal control (NC), MS, MS with metformin 200 mg/bw and simvastatin 10 mg/kg body weight (DMS), 100 and 200/bw GCE (GCE 100 and GCE 200, respectively). The extract was given through oral gavage daily. Extract dose was given in milliliters based on the body weight measured weekly. The food and water intake was recorded daily. At the end of the experimental period, the animals were anesthetized with ether following a 12 hrs fasting period. Blood samples were drawn from the heart into microcentrifuge tube, and serum samples were obtained by centrifugation at 4000 ×g for 15 minutes at 4°C. All animal experiments protocol were reviewed and approved by the Ethics Committee of Faculty of Medicine, Brawijaya University.

Physiological measurement
Daily food intake and fluid intake were measured every day, and body weight was measured every week. The food and fluid intake for each rat were measured by subtracting the provided amount by the remaining amounts in the cage.

Biochemical analysis
The serum concentrations of TG, HDL-cholesterol, and fasting glucose were measured enzymatically using commercial kits (Biolabo, France). Serum adiponectin and insulin levels were analyzed using enzyme-linked immunosorbent assay method (ElabsScience, China). The HOMA-IR was used to calculate an index from the product of the fasting concentrations of plasma glucose (mg/dL) and insulin (µu/L) divided by 4.1. Lower HOMA-IR values indicate greater insulin sensitivity and higher HOMA-IR values indicate IR.

Blood pressure measurements
Blood pressure was measured using the tail-cuff method with sphygmonanometer technique at the baseline and at the end of the experiment. Three readings were taken consecutively, and the average was then calculated and taken as a final reading for SBP.

Statistical analysis
All data were analyzed with Statistical Package for Social Sciences (SPSS, version 22) and were presented as mean values with their standard deviation and subjected to one-way ANOVA, independent t-test, and pair t-test with significant p<0.05.

RESULT
Effect of GCE on changes in the body weight, food, and water intake
The body weight, food, and water intake of experimental rats were elevated during 7 weeks. However, at the end of study, no significant difference was observed in those variables among all groups as presented in Tables 1 and 2, respectively. Body weight increased significantly in 200 GCE group during 7-week intervention. Moreover, a significant food intake increase was observed in DMS group. Furthermore, a significant water intake increase was observed in all experimental groups during 7 weeks.

Effects of GCE on blood biochemistry
Effect of GCE on fasting blood glucose, TG, and HDL cholesterol on the experimental animals were presented in Tables 3 and 4. The level of fasting blood glucose TG and HDL was not statistically different among all groups at the baseline. Furthermore, after intervention for 7 weeks, a significant difference on fasting blood glucose, and TG level was observed in all interventional groups (p<0.05) compared to that of the baseline. At the end of study, a significant difference on fasting blood glucose and TG level was observed in GCE 100, GCE 200, and DMS group compared to that of MS group. The lowest fasting bloodefasting glucose level was observed in DMS and GCE 100 groups respectively. Moreover, the lowest TG level was observed in DMS and GCE 200 group, respectively. No statistically different observed on HDL level among all interventional group.

Effect green coffee on blood pressure
GCE 100 and GCE 200 group showed a significant SBP decrease after intervention for seven weeks compared to that of the baseline. At the end of study, a significant lower SBP was observed in GCE 100 and GCE 200 group compared to that of MS group (Table 4).

Effects of GCE on plasma adiponectin
Effect of GCE on plasma adiponectin level was presented in Fig. 1. GCE 100 and GCE 200 group showed significantly higher adiponectin level compared to that of MS and DMS group. Moreover, administration of GCE at a dose of 100 mg/kg bw resulted in a slightly higher adiponectin level compared to that of 200 mg/kg bw.

Effects of GCE on HOMA-IR index
Effect of GCE on HOMA-IR index was presented in Fig. 2. GCE 100, GCE 200, and DMS group showed significantly lower HOMA-IR index compared to that of MS group. Moreover, GCE 100 group showed lower HOMA-IR index compared to that of GCE 200 group although it was not achieve any significant statistical difference. At the end of study, a significant different of HOMA-IR index was observed in GCE 100 and GCE 200 group compared to that of MS group (p<0.05).

DISCUSSION
A recent study was conducted in rat model that met the criteria of MS (hyperglycemia, elevated TG level, decreased HDL level, and

| Table 1: Effect of GCE on changes in the body weight of normal and MS rats |
|-------------|-----|-----|
| Experimental groups | Body weight (g) |
|               | Initial | Final |
| NC           | 297.8±10.66 | 411±9.3  |
| MS           | 336.4±14.7  | 386.4±57.17 |
| GCE 100      | 298.4±39.87 | 311±59.74 |
| GCE 200      | 301±44.32    | 351.6±74* |
| DMS          | 310±30.05    | 318±44.07 |

Values are mean±SD, n=5. *Significant between baseline and after 7 week (p<0.05). NC: Normal control, MS: Metabolic syndrome, GCE: Green coffee extract.
Table 2: Effect of GCE on changes in food intake and fluid intake of normal and metabolic syndrome rats

<table>
<thead>
<tr>
<th>Experimental groups</th>
<th>Food intake (g)</th>
<th>Fluid intake (ml)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Initial</td>
<td>Final</td>
</tr>
<tr>
<td>NC</td>
<td>21.8±1.49</td>
<td>26.85±1.9</td>
</tr>
<tr>
<td>MS</td>
<td>16.8±0.44</td>
<td>45.4±14.02</td>
</tr>
<tr>
<td>GCE 100</td>
<td>18.2±1.48</td>
<td>45.2±4.88</td>
</tr>
<tr>
<td>GCE 200</td>
<td>19.38±2.58</td>
<td>42.97±13.46</td>
</tr>
<tr>
<td>DMS</td>
<td>18.87±2.54</td>
<td>41.81±9.02</td>
</tr>
</tbody>
</table>

Values are mean±SD, n=5. *Significant between baseline and after 7 week (p<0.05). NC: Normal control, MS: Metabolic syndrome, GCE: Green coffee extract, SD: Standard deviation

Table 3: Effect of GCE on changes in fasting blood glucose and TG of normal and metabolic syndrome rats

<table>
<thead>
<tr>
<th>Experimental groups</th>
<th>Fasting blood glucose (mg/dl)</th>
<th>TG (mg/dl)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Initial</td>
<td>Final</td>
</tr>
<tr>
<td>NC</td>
<td>92.2±6.84</td>
<td>95.2±11.45</td>
</tr>
<tr>
<td>MS</td>
<td>255±29.72</td>
<td>270.2±40.18</td>
</tr>
<tr>
<td>GCE 100</td>
<td>260±26.16</td>
<td>186.20±11.10**</td>
</tr>
<tr>
<td>GCE 200</td>
<td>247.80±29.21</td>
<td>201.20±11.3**</td>
</tr>
<tr>
<td>DMS</td>
<td>248±28.67</td>
<td>178.40±59.94**</td>
</tr>
</tbody>
</table>

Values are mean±SD, n=5. *Significant between baseline and after 7 week (p<0.05). *Significant compared to that of MS group (p<0.05). NC: Normal control, MS: Metabolic syndrome, TG: Triglyceride, SD: Standard deviation, GCE: Green coffee extract

Table 4: Effect of GCE on changes in HDL-cholesterol and SBP of normal and MS rats

<table>
<thead>
<tr>
<th>Experimental groups</th>
<th>HDL-cholesterol (mg/dl)</th>
<th>SBP (mm/Hg)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Initial</td>
<td>Final</td>
</tr>
<tr>
<td>NC</td>
<td>44.40±3.13</td>
<td>44.80±3.03</td>
</tr>
<tr>
<td>MS</td>
<td>32.20±6.76</td>
<td>34±3.39</td>
</tr>
<tr>
<td>GCE 100</td>
<td>27.6±8.17</td>
<td>40.40±5.77*</td>
</tr>
<tr>
<td>GCE 200</td>
<td>34.60±5.45</td>
<td>39.20±2.86</td>
</tr>
<tr>
<td>DMS</td>
<td>29±8.09</td>
<td>43±7.07</td>
</tr>
</tbody>
</table>

Values are mean±SD, n=5. *Significant between baseline and after 7 weeks (p<0.05). *Significant compared to that of MS group (p<0.05). NC: Normal control, MS: Metabolic syndrome, SBP: Systolic blood pressure, SD: Standard deviation, HDL: High-density lipoprotein, GCE: Green coffee extract

Adiponectin is a key molecule in the pathogenesis of MS [19]. In the present study, GCE improved metabolic profile of MS through adiponectin increase [20,21]. This study demonstrated a significantly higher adiponectin level in GCE 100 and GCE 200 compared to that of MS group. The adiponectin level also closed to that of NC group. This study was comparable with the previous study by Cho et al that revealed a significantly higher adiponectin level after 8 weeks administration of 0.02% CGA in diet compared to that of obese mice group [20]. Moreover, a study by Ong et al suggested that 2 weeks treatment of CGA from green coffee bean increased serum adiponectin level in Lepr+/- mice compared to that of diabetic mice [10]. To the extent of our knowledge, this study was the first study that proved the benefit of GCE in MS rat.

Adiponectin level increasing effect by GCE was mediated by CGA induce transcriptional activity of peroxisome proliferator-activated receptor gamma (PPAR-γ). The previous study by Wu et al showed that 200 and 400 mg of CGA for 12 weeks increased transcriptional activity of PPAR-γ and consequently adiponectin level in ApoE-/- mice [22]. It was revealed that PPAR-γ affected adiponectin by controlling the expression of the adiponectin genes in adipose tissue. Moreover, adiponectin increasing effect was also shown in DMS group compared to that of MS group. Nevertheless, the effect of metformin and simvastatin only restoring the adiponectin level as high as NC group's level. This result was consistent with the previous study that suggested metformin effect on restoring the adiponectin level as high as normal rat [23]. Thus, GCE administration demonstrated a better adiponectin increasing effect compared to that of 200 mg/kg worm metformin and 10 mg/kg worm simvastatin administration.

This study revealed that adiponectin level of GCE 100 was higher than that of GCE 200 although it was not achieve any statistical significance. In contrast, Choi et al suggested that higher GCE dose resulted in higher adiponectin level [21]. The differences of substances composition
in the extract might contribute to this result. The extract used in our study was not purified, some caffeine and other polyphenols might contain in our extract on the other hand, Choi et al. used GCE contained 50% of caffeoylquinic acid (CGA) [21]. The chronic effect of higher GCE dose administration on homocysteine level might explain lower adiponectin level in higher GCE dose. The previous studies demonstrated that chronic administration of high-dose green coffee raised blood homocysteine level [24]. The mechanisms by which CGA raise plasma homocysteine are not clear. The previous study suggested that caffeine content in GCE might contribute significantly to the high level of homocysteine. The previous study revealed that pure caffeine consumption increased homocysteine concentrations in healthy subjects [25]. However, caffeine from paper-filtered coffee only contributed to 25-50% of homocysteine increase compared to that of a similar amount of caffeine [25].

High homocysteine level has been proved as a suppressor of adiponectin production. The previous study suggested that liver might affect adipocyte physiology through adipokines expression regulation through the secretion of metabolic molecules such as homocysteine [27]. The previous study clearly demonstrated that multiple pathways were involved in this mechanism. Homocysteine induced both inhibition of adiponectin gene expression in primary adipocytes and reduced circulating adiponectin levels in an animal model of mild hyperhomocysteinemia [27]. It revealed that elevated homocysteine in adipose tissue may play a causal role in suppressing adiponectin production.

Adiponectin also known as insulin sensitizer agent. The present study suggested that higher adiponectin negatively correlated with HOMA-IR index [14]. In our study, we assumed that higher adiponectin level led to lower HOMA-IR index. GCE 100 group who had higher adiponectin level showed lower HOMA-IR index compared to that of GCE 200 and DMS group. Although there was no significant, HOMA-IR difference among GCE 100, GCE 200, and DMS. The obtained HOMA-IR index by GCE 100 closed to that of NC group. These findings are consistent with previous reports that adiponectin functions as an insulin sensitizer by decreasing hepatic glucose output and thereby contributing to the regulation of whole-body glucose homeostasis [28]. Consistent with this, restoration of adiponectin levels in vivo by CGA from green coffee showed improvement of glycemia, insulin sensitivity, and fatty acid metabolism [29]. Moreover, it could have enhanced the activation of adenosine monophosphate-activated protein kinase (AMPK) through calmodulin-dependent protein kinase in muscle and liver [10] and upregulated AdipoR2 expression that resulted in insulin sensitivity modulation by stimulation of glucose utilization and fatty acid oxidation through phosphorylation [30]. These mechanisms were considered as beneficial effect of GCE in metabolic alterations.

Recently, this study showed that 100 mg/kg bw GCE was the effective dose in treating hyperglycemia, dyslipidemia in rat model of MS for 7 weeks. The extent of GCE 100 mg/kg bw decreased fasting blood glucose, and TG level was also shown in GCE 200 mg/kg bw. However, the fasting blood glucose and TG level were slightly lower than GCE 100 mg/kg bw. In contrast, the previous study by Shimoda showed that an oral administration of GCE (200 and 400 mg/kg/day) for 13 days resulted in a tendency of serum TG level reduction in mice [31]. Moreover, Choi et al. reported that an oral administration of GCE 100 and GCE 200 in C57BL mice reduced serum glucose, TG level after 6 weeks, and significantly difference compare to that of HFD group [21]. On the other hand, in significant serum HDL increase in GCE 100 and GCE 200 group was observed in this study. Antihyperglycemia and antidiyslipidemia of GCE might be mediated by CGA activity. CGA affected AMPK signaling pathway [32], upregulated expression of adiponectin [10], upregulated adiponectin receptor gene [30], and increased transcriptional factor of PPAR-α [20] and PPAR-γ [21] which could regulate glucose and lipid metabolism.

A recent study revealed no body weight lowering effect after GCE administration for 7 weeks using both doses. This result was consistent with a study by Panchal et al. that showed no body weight lowering effect of 5% aqueous coffee extract oral administration for 8 weeks in MS rat model induced by HFD and high carbohydrate [33]. In contrast, other study revealed that GCE administration resulted in a tendency of visceral fat and body weight reduction. Song et al. suggested that the administration of 0.3% decaffeinated GCE for 11 weeks in HFD-fed mice resulted in body weight reduction [13]. Shimoda et al. also suggested that the administration of 0.5% and 1% GCE reduced visceral fat content and body weight.

The present study shows that GCE 100 and 200 mg/kg bw reduced blood pressure from 152.6±3.84 to 145.4±3.43 and 152.4±7.40 to 147.0±8.03, respectively. However, blood pressure increased was observed in MS group and DMS group. The blood pressure lowering effect of GCE was first reported in 2002 in spontaneous hypertensive rat (SHR) rat [34]. The previous study by Panchal et al. demonstrated that oral administration of 5% aqueous coffee extract after 8 weeks. SBP were reduced by 10-15 mmHg from baseline (measured by tail cuff), and the effect persisted through the end of the experiment [33]. As mentioned previously, GCE are rich in CGAs, with the principal form being 5-CQA. The presence of 5-CQA was hypothesized to promote SBP reduction. Both acute and chronic administration of 5-CQA showed almost identical results on SBP as seen in GCE administration on MS animal model diet induced. Furthermore, several mechanisms of SBP reduction by CGA and it metabolites such as, inhibition of nicotinamide adenine dinucleotide phosphate oxidase expression and activity, reduced free radical production, directly scavenging free radicals, stimulation of NO production by the endothelial-dependent pathway, and inhibition of angiotensin-converting enzyme in the plasma [35]. Hypotension effect of GCE could be influenced by roasting condition which that increase antioxidant activity [36].

Roasting effect on coffee can increase the antioxidant effect of polyphenols contained in coffee. Castillo et al. analyzed various roasted degree of coffee beans by gel filtration chromatography, ultraviolet- visible spectrophotometry, capillary electrophoresis, and the 2,2'-azino-bis(3-ethylbenzthiazoline-6-sulfonic acid) assay. It revealed that maximum antioxidant activity was observed in light to medium roasted coffee; unroasted and dark coffee had significantly lower antioxidant activity [36].

Our study regarding these issues is warranted.
CONCLUSION

This study revealed the beneficial effect of light-roasted GCE administration on the improvement of adiponectin level, HOMA-IR index, lipid profile, fasting blood glucose level, and SBP in MS rat model.

ACKNOWLEDGMENT

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