Correlation Between Adiponectin Receptor with Indices of Glucose Homeostasis and Mediators Of Insulin Sensitivity in Type 2 Diabetic Rats Treated With Puguntano (Curanga felterrae Lour.) Leaf Extract

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ABSTRACT. Objective: Adiponectin receptors (AdipoR) regulates metabolism and has anti-inflammatory and insulin-sensitizing effects. We aimed to determine the relationships between AdipoR with parameters of glucose homeostasis (FPG, insulin, and HOMA-IR), and insulin sensitivity (PPAR-γ and p38MAPK) in T2DM rats treated with puguntano (Curanga felterrae Lour.) leaf extract. Methods: T2DM was induced in Wistar rats aged 8–10 weeks and weighing 180–200 g by high-fat diet (HFD) feeding and low-dose streptozotocin (30mg/kg.bw) administration. The rats were then allocated randomly to a treatment group and a control group (n=24 each). The treatment group was orally administered puguntano leaf extract (200 mg/kg.bb) once daily for 10 days. Subsequently, FPG and plasma insulin were measured, and HOMA-IR was calculated. Results: There was significantly difference between treatment group and control group on AdipoR and parameter of glucose homeostasis (FPG, insulin, and HOMA-IR) and insulin sensitivity (PPAR-γ, p38MAPK (all, p<0.01). In both groups, there were no significantly correlations between of AdipoR with all parameter of glucose homeostasis and insulin sensitivity except PPAR-γ (p <0.003) across the entire cohort of rats. Conclusion: Our data suggest that puguntano could improve glucose homeostasis, insulin sensitivity and molecular mediators of insulin sensitivity. There were no significantly correlated between improvement of AdipoR with glucose homeostasis and molecular mediators of insulin sensitivity in T2DM.

Keywords: AdipoR, Type 2 Diabetes Mellitus, glucose homeostasis, Insulin sensitivity, Curanga fel-terrae Lour.

1 INTRODUCTION

Adiponectin is an insulin-sensitizing adipokine that reduces the plasma glucose concentration of rats by suppressing liver glucose production and increasing peripheral glucose uptake, independent of insulin [Berg et al., 2001]. The mechanisms of these effects involve the downregulation of hepatic phosphoenolpyruvate carboxykinase and glucose-6-phosphatase expression [Yamauchi et al., 2002], and the activation of AMP-activated protein kinase (AMPK), which ameliorates insulin resistance and prevents hepatosteatosis [Liu et al., 2012].

Adiponectin has also been shown to increase fatty acid oxidation in skeletal muscle in vivo, thereby reducing triglyceride storage in muscle and liver, and increasing insulin sensitivity [Yamauchi et al., 2001]. Furthermore, adiponectin suppresses inflammation associated with macrophage infiltration into insulin target tissues [Iannitti et al., 2015].

The effects of adiponectin are mediated by binding to the adiponectin receptors (AdipoR1 and AdipoR2), and regulation of the expression of metabolic genes and insulin sensitivity in insulin target tissues [Yamauchi et al., 2007]. Higher levels of expression of both AdipoRs ameliorates insulin resistance, modulates food intake and energy expenditure, and reduces inflammation. Furthermore, the mRNA expression levels of the AdipoRs correlate with adiponectin concentrations [Yamauchi et al., 2007; Yamauchi and Kadowaki, 2008].

Previous studies have shown that administration of an ethanolic extract of puguntano leaf (*Curanga felterrae* Lour.) significantly improves glucose metabolism, ameliorates insulin resistance, increases adiponectin concentration [Lindarto et al., 2016], and increases the expression of AdipoR [Lindarto et al., 2019], p38 mitogen-activated protein kinase [p38MAPK], and glucose transporter-4 [GLUT-4] (Syafiril et al., 2019) in type 2 diabetes mellitus (T2DM) rats.

We aimed to determine the relationships between AdipoR with indices of glucose homeostasis: fasting plasma glucose (FPG), insulin, homeostasis model assessment-insulin resistance (HOMA-IR); and molecular mediators of insulin sensitivity: peroxisome proliferator-activated receptor-γ (PPAR-γ) and p38MAPK in T2DM rats treated with puguntano (*Curanga felterrae* Lour.) leaf extract.
2 METHODS

We studied 48 male Wistar rats aged 8–10 weeks and weighing 180–200 g that were maintained under a natural light cycle at a temperature of 22–25°C. T2DM was induced by feeding a high-fat diet (HFD) for 5 weeks, followed by the intraperitoneal injection of 30 mg/kg streptozotocin [Sigma-Aldrich, Munich, Germany].

The control group was sacrificed when diagnosed with T2DM and the treatment group was sacrificed after the completion of puguntano treatment. This was accomplished by the induction of anesthesia using ketamine, followed by decapitation.

Thereafter, blood was then collected from the left ventricle, and FPG (spectrophotometry) was used to measure glucose concentration, and diabetes was considered to be present when the FPG was > 200 mg/dL [Zhang et al, 2008], and fasting insulin (sandwich ELISA) were measured. Skeletal muscles were dissected and then homogenized in a cold homogenization buffer (−80°C). The protein levels of p38-MAPK, PPAR-γ, and AdipoR were determined in these homogenates using a Qayeebio kit (China). The rats were then allocated at random to a treatment group and a control group (n=24 each).

The ethanolic extract of puguntano leaves was obtained by maceration in the Department of Biological Pharmacy, Faculty of Pharmacy, Universitas Sumatera Utara, Medan, Indonesia. The treatment group was administered an ethanolic extract of puguntano leaf in carboxymethylcellulose-Na (CMC Na; 0.5% solution; 200 mg/kg/day) by gavage daily for 10 days [Ministry of Health Republic of Indonesia, 2013].

This research has been approved by the Ethics Committee of Faculty Medicine and H. Adam Malik Hospital (Reference 42 /TGL/ KPEK FK USU-RSUP HAM/2018).

Statistical Analysis

Statistical analysis was performed using SPSS 22.0 [IBM, Inc., Armonk, NY, USA]. All data are expressed as means ± standard deviations. The Wilcoxon test was used to compare non-normally distributed data, and Pearson’s or Spearman’s correlations were calculated for pairs of variables. A p-value < 0.05 was taken to indicate statistical significance.

3 RESULTS AND DISCUSSION

There were significant differences in FPG, plasma insulin, HOMA-IR, and muscle PPAR-γ, p38 MAPK, and AdipoR protein levels between the treatment and control groups (Table 1).
Table 1  Comparison between treatment and control groups on parameter of glucose homeostasis and insulin sensitivity

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Treatment group (n=24)</th>
<th>Control group (n=24)</th>
<th>p*</th>
</tr>
</thead>
<tbody>
<tr>
<td>FPG (mg/dl)</td>
<td>136.63±33.62</td>
<td>375.58±29.15</td>
<td>0.001**</td>
</tr>
<tr>
<td>Insulin (nU/L)</td>
<td>52.32±3.32</td>
<td>57.36±6.28</td>
<td>0.001**</td>
</tr>
<tr>
<td>HOMA-IR</td>
<td>0.86±0.20</td>
<td>3.05±0.51</td>
<td>0.001**</td>
</tr>
<tr>
<td>PPAR-γ (ng/mL)</td>
<td>40.80±6.83</td>
<td>29.56±1.06</td>
<td>0.001**</td>
</tr>
<tr>
<td>p38MAPK (ng/mL)</td>
<td>23.70±4.04</td>
<td>20.81±3.02</td>
<td>0.005**</td>
</tr>
<tr>
<td>AdipoR (ng/mL)</td>
<td>16.64±3.83</td>
<td>13.79±1.47</td>
<td>0.001**</td>
</tr>
</tbody>
</table>

Data are expressed as mean ± standard deviation. *The Wilcoxon test was used to evaluate differences between the control and treatment groups. FPG: fasting plasma glucose; HOMA-IR: homeostatic model assessment-insulin resistance; PPAR-γ: peroxisome proliferator-activated receptor-γ; p38 MAPK: p38 mitogen-activated protein kinase

Total muscle AdipoR protein levels significantly correlated with FPG, insulin, HOMA-IR, and muscle PPAR-γ protein across all the rats studied. However, there were no correlations identified separately in the control and treatment groups, except for one between total muscle AdipoR protein and muscle p38MAPK protein in the treatment group (Table 2).

Table 2  Correlations of AdipoR with indices of glucose homeostasis and molecular mediators of insulin sensitivity in each group and across all the rats studied.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Treatment Group (n=24)</th>
<th>Control group (n=24)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>r          p</td>
<td>r          p</td>
</tr>
<tr>
<td>FPG (mg/dl)</td>
<td>−0.264 0.212</td>
<td>−0.27 0.196</td>
</tr>
<tr>
<td>Insulin (nU/L)</td>
<td>−0.402 0.052</td>
<td>0.024 0.912</td>
</tr>
<tr>
<td>HOMA-IR</td>
<td>−0.254 0.231</td>
<td>−0.03 0.878</td>
</tr>
<tr>
<td>PPAR-γ (ng/mL)</td>
<td>0.578 ** 0.003</td>
<td>0.088 0.680</td>
</tr>
<tr>
<td>p38MAPK (ng/mL)</td>
<td>−0.273 0.197</td>
<td>−0.33 0.106</td>
</tr>
</tbody>
</table>

Data are expressed as means ± standard deviations. FPG: fasting plasma glucose; HOMA-IR: homeostatic model assessment-insulin resistance; PPAR-γ: peroxisome proliferator-activated receptor-γ; p38 MAPK: p38 mitogen-activated protein kinase. Statistically significant correlations are shown in bold
4 Discussion

The secondary metabolites derived from ethanolic extracts of puguntano leaf have been identified to be glycosides [Zhou, 2005], flavonoids [Huang, 1998], saponins [Fang et al, 2009], terpenoids [Wang et al, 2006], and these have been shown to reduce blood glucose by stimulating the synthesis and secretion of insulin. In addition, tannins increase glucose uptake by activating the phosphoinositide 3-kinase (PI3K) and p38MAPK signaling pathways and promoting GLUT-4 translocation [Kumari and Tannins, 2012].

AdipoRs mediate improvements in glucose metabolism through mechanisms such as an increase in muscle fatty acid oxidation and the suppression of lipid accumulation in the muscle and liver, which improves insulin sensitivity. These effects are in large part thought to be mediated through the activation of AMPK [Yamauchi et al, 2001]. However, adiponectin also activates the p38MAPK pathway [Mao et al, 2006], which has anti-inflammatory effects [Xin et al, 2011], and increases PPAR-γ expression, thus promoting adipocyte differentiation [Fu et al, 2005]. Furthermore, in diabetic patients, adiponectin concentrations demonstrate a significant negative correlation with body mass index and positive correlations with systolic blood pressure and microalbuminuria [El Dayem et al, 2015]. Treatment with a combination of 0.2 mg/ml of a water/ethanol extract of *Momordica charantia* fruit and seeds and 0.5 nM insulin significantly increases glucose uptake and adiponectin secretion in 3T3-L1 adipocytes [Roffey et al, 2007]. In addition, administration of an American ginseng (*Panax quinquefolius*) extract containing a quantifiable amount of ginsenosides reduces cell growth and lipid accumulation, and increases adiponectin expression in 3T3-L1 cells [Yeo et al, 2011]. Adiponectin also has anti-atherogenic effects and its effects are similar, but additive to, the effects of insulin on metabolism and the vascular endothelium. However, the best-known relationships are the inverse relationships between adiponectin concentration and obesity and insulin resistance [Balsan et al, 2015]. Finally, AdipoR ameliorates diabetes associated with obesity and increases exercise endurance, which prolongs the shortened lifespan of obese mice fed an HFD.

We don’t have identified significant correlations between muscle AdipoR proteins, indicators of glucose homeostasis, and key molecular mediators of insulin sensitivity except PPAR-γ in muscle in T2DM rats. It is expected that the current findings will contribute to the elucidation of AdipoR-mediated signal transduction and further encourage the development and optimization of AdipoR-targeting therapeutics for obesity-related diseases, such as diabetes [Okada-Iwabu et al, 2015].

5 Conclusion

The study suggest that puguntano could improve glucose homeostasis, insulin sensitivity and molecular mediators of insulin sensitivity. There were no significantly correlated between
improvement of AdipoR with glucose homeostasis and molecular mediators of insulin sensitivity in T2DM.

REFERENCES


