Evaluation of PRKAA2 Genetic Variation on Metformin Efficacy as an Initial Therapy Among Drug-Naïve Patients With Type II Diabetes Mellitus

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Background: Metformin is the most popular oral antidiabetic agent, which is recommended as initial monotherapy. AMPK is the pivotal target of metformin molecular mechanisms. AMPK subunit a2 (encoded by PRKAA2) is a gene contributable to increase type 2 diabetes mellitus (T2DM) risk. This study aimed to evaluate PRKAA2 rs2796498, rs2746342, and rs980799 genetic variations on metformin efficacy.

Methods: This study enrolled 191 newly diagnosed Indonesia T2DM patients in primary health care. Patients who received metformin as monotherapy for at least 3 months were included for genotyping. Genotyping was performed using the Taqman assay.

Results: Baseline characteristics showed that BMI was higher among AA than GG+AG (p=0.04). Patients with TT genotype showed a higher FBG and HbA1c than GG+GT (p=0.02 and p=0.02, respectively). There was no significant difference in allele and genotype frequencies between responders and non-responders group in PRKAA2 rs2796498, rs9803799, and rs2746342. However, among PRKAA2 rs2796498, AG carrier had 0.32 times of responding in metformin efficacy after adjusting BMI, WC, blood pressure, lipid profiles, and eGFR. Dominant model of rs2796498 showed a significant association (OR=0.34, 95%CI=0.13 – 0.90) to metformin efficacy.

Conclusions: Our findings suggest that PRKAA2 rs2796498 genetic variation may affect metformin efficacy, especially AG carrier, in drug-naïve T2DM patients.

Introduction
Metformin is notable as the most prescribed medicine for T2DM patients as a recommended initial type 2 diabetes mellitus (T2DM) therapy by many international guidelines. Drug-naïve T2DM patients who have HbA1c 7-9% are suggested to consume metformin until it reaches the maximum doses before adding a second agent [1, 2]. Metformin considerably has better safety and efficacy as initial T2DM therapy [3]. Metformin is a guanidine derivate which acts as an unchanged form. Metformin specifically reduces blood glucose by enhancing glucose uptake in peripheral tissue and suppressing glucose hepatic production [4–6]. The underlying complex molecular mechanism of metformin has been observed in many studies, and AMP-activated protein kinase (AMPK) was mentioned to have a pivotal role in metformin action [7, 8].

AMPK is a cellular energy sensor whose phosphorylation is stimulated by lower ATP concentration in cells. AMPK has three subunits consisting of a, b, and g. AMPK suppresses the anabolic process, which utilizes ATP [9]. Metformin stimulates phosphorylation of Thr-172 at a catalytic unit [10]. Some studies also demonstrated that metformin activates AMPK after inhibiting mitochondrial complex I respiratory chain. Thus it reduces the ATP level [8, 11]. Furthermore, other studies reported that LKB-1 is also involved in AMPK activation by metformin [12]. Besides, AMPK via acetyl-CoA carboxylase 1 (ACC1) and ACC2 could inhibit malonyl CoA synthesis. As a result, it reduces lipogenesis and increases insulin sensitivity [13]. Therefore, metformin does not affect weight gain.

Nevertheless, there is a different efficacy of metformin among T2DM patients [14]. Metformin efficacy is affected by various factors, including age, lifestyle, baseline HbA1c, adherence [15, 16], gene-drug interaction, and gene-environment interaction [17]. Inter-individual variability plays an important role in drug response affected by drug-gene interaction. Therefore, pharmacogenomic still be a promising study for discovering the relationship between gene and medicine's pharmacodynamic and pharmacokinetic, in this context especially metformin efficacy.

Many studies have explored the effect of genetic variation on metformin response in the pharmacokinetic area [18, 19]. Otherwise, a little study conduct pharmacogenomic research focused on the pharmacodynamic of metformin. Accordingly, it is relevant to explore pharmacogenetic concerning the pharmacodynamic of metformin in achieving the HbA1c goal. Since AMPK is involved as the main target in the metformin molecular mechanism, genes coding AMPK may have contributable to metformin efficacy. A recent review declared that PRKAA2 coding AMPKa2 is one of a gene related to clinical outcomes after metformin therapy [20]. Nevertheless, that review has not mentioned an SNP of PRKAA2 contributing to metformin response yet.

Previous studies have reported an association of PRKAA2 with susceptibility of T2DM (rs2796498 and rs2746342) [21, 22], even in metformin effectiveness (rs9803799) [23]. However, no studies observe the association between PRKAA2 rs2796498 and rs2746342 genetic variation and the effectiveness of metformin use. Notably, those three SNP have not been explored yet in Indonesia. Therefore, this study aimed to observe the genotype frequency among T2DM Indonesian patients receiving metformin. Furthermore, this study investigates the influence of PRKAA2 rs2796498, rs9803799, and rs2746342 genetic variation on the efficacy of metformin in drug-naïve T2DM patients, determined by the reduction of HbA1c level.

Methods
Participants
This study recruited 191 participants from ten primary health care (PHC) in Yogyakarta Province, Indonesia. Physicians in PHC gave the first prescription of metformin on newly diagnosed T2DM patients. Then participants were followed up for three months. The participant's selection following inclusion criteria: newly diagnosed T2DM patients with HbA1c>7%, 18-70 years old, do not consume other hyperglycemia agents, either oral or injectable dosage forms. Participants were excluded when GFR<30 ml/min/m2, albumin serum 3.4 – 4.8 g/dL, and diagnosed as diabetes gestational. Finally, only 106 newly diagnosed T2DM patients who had taken the first prescription metformin as monotherapy and completed follow up for three consecutive months were enrolled in this study. Patients were classified into two groups: metformin responders (n=45) and metformin non-responders (n=61). All of our participants were of Indonesia origin. Metformin responders are patients who could achieve decreasing HbA1c>1.12% during follow up [24].

Ethics
Anthropometric measurement, blood pressure, blood sample collecting for clinical chemistry and genotyping analysis, and medical record data were obtained for this study after patients signing informed consent. All participants were briefed about the study aim, procedure, duration, potential risk, and benefit. The study was performed in compliance, according to the Declaration of Helsinki. The study protocol was approved by the Medical and Health Research Ethics Committee (MHREC) Faculty of Medicine, Public Health, and Nursing Universitas Gadjah Mada – Dr. Sardjito General Hospital. Patients had the right not to participate in our study at any time.

Clinical measurement

Our study collected demographic data, anthropometric data, and laboratory results. Age and gender, as demographic data and metformin dose, were obtained from medical records. Nutritionists measured anthropometric measurements, including body weight, height, and waist circumference. Body Mass Index (BMI) was calculated by divided body weight (kg) by height (m²). Blood pressure was measured by the nurse in early registration. We examined laboratory results, including FBG, HbA1c, creatinine serum, HDL-c, Triglyceride-c, and total cholesterol. Laboratory data were collected by a PHC analyst that helped by a commercial laboratory after an overnight fast. FBG and creatinine serum was measured by the hexokinase method and enzymatic method, respectively. HbA1c was calculated using high-performance liquid chromatography (Cobas D-10). Glomerular filtration rate (GFR) was calculated using CKD-EPI formulation using the creatinine level. Lipid profiles, including HDL-c, Triglyceride-c, and total cholesterol, were measured using Cobas C311. LDL was calculated using Friedewald formulation.

DNA extraction and genotyping

According to the kit protocol, genomic DNA was extracted from peripheral whole blood-EDTA using Geneaid® Blood DNA Mini Kit and stored at -20 until the genotyping procedure. Genotyping in rs2796498, rs9803799, and rs2746342 was performed using the TaqMan® genotyping assay and Applied Biosystems® qPCR 7500 Fast Real-Time PCR System. The final reaction volume using in real-time PCR is 10mL, including 2.5 mL nucleotide-free water, 5mL TaqMan GTXpress mix, 0.5mL TaqMan SNP genotyping assay, and 2mL of genomic DNA. The thermal cycle for a reaction was as follows: 40 cycles at hold 95°C for 20s, at denaturing 95°C for 3s, and then annealing 60°C for 30s. PRKAA2 SNPs were determined using the following primer sequences:

rs2796498: CTGTAACGTTATGCTTTAACAGTGA [A/G] GAGAGCAACCTTACCTTTTGCTAG
rs9803799: TAAATACAGGGTTTATATCCCCACA [G/T] TCAATGTAAATTCCTTTTTAAAA
rs2746342: AGAGAGGCTAAGATGCAGGCTGTAC [G/T] CTGGGTAGCCATGTACTCAGTTGTA

Statistical analysis

The baseline and follow-up participant's characteristics were analyzed and compared between the genotype model using an independent t-test or chi-square, as appropriate. We applied the dominant and recessive model in comparing clinical characteristics. Data are expressed in mean±SD for numerical data and n(%) for categorical data. Allele and genotype frequencies were evaluated by Hardy-Weinberg equilibrium (HWE) using chi-square. The association between responder status and genotypes was analyzed using multinominal logistic regression by adjusting for age, gender, BMI, WC, lipid profiles, glomerular filtration rate, and blood pressure. All statistical analysis was performed using SPSS version 25.0, and p<0.05 was considered statistically significant.

Results

Of the 191 participants who have classical T2DM signs, 62 were dropped out: 35 were HbA1c<7%, 3 were age>70 years old, 18 had eGFR<30 mL/min/m², 20 did not comply with consuming metformin, and 23 needed the second agent. The baseline characteristics are shown in Table. 1. BMI in the AA group (29.00±5.34) was higher than the GG+AG group (25.01±4.21) in the rs2796498 recessive model (p=0.04). However, both have been classified as obesity. The differences between FBG and HbA1c before metformin treatment are only found in the rs2796498 recessive model (p=0.02 and p=0.04, respectively). Specifically, rs2796498 in the recessive model showed that the GG+GT group tended to have lower FBG and HbA1c before metformin therapy (174.39±58.08 mg/dL and 9.29±1.74%) than the TT group (211.67±74.21 mg/dL and 10.27±2.07%). Moreover, there were no significant differences in baseline characteristics in rs9803797, either dominant or recessive models.

Table 1. Baseline characteristics in drug-naïve T2DM patients before receiving monotherapy metformin based on PRKAA2 genetic variation
### Allele and genotype frequency of PRKAA2 genetic variation in responders and non-responder of metformin therapy

Allele and genotype frequency of PRKAA2 genetic variation in responders and non-responder of metformin therapy are listed in Table 2. The HWE of rs2796498, rs9803799, and rs2746342 were 0.47, 0.03, and 0.92, respectively. It suggested that the PRKAA2 genotypes of rs2796498 and rs2746342 in our population are consistent with HWE, but not rs9803799 (p<0.05). Minor allele frequencies (MAFs) of rs2796498, rs9803799, and rs2746342 were 27.8% and 22.1%, 10.0% and 4.9%, 44.4% and 39.3%, respectively, in responders and non-responders group. As presented in Table 2., the frequency of allele G, T, and G of rs2796498, rs9803799, and rs2746342, respectively, tend to higher in metformin non-responders than responders group. Nonetheless, we could not find the difference statistically of allele frequency between responders and non-responders (p=0.35 for rs2796498, p=0.15 for rs9803799, and p=0.46 for rs2746342).

We could not find any difference in blood pressure, FBG, HbA1c, HBA1c change, and lipid profiles after receiving metformin therapy based on PRKAA2 genetic variation (Table. 3).

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>rs2796498</th>
<th>rs9803799</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Dominant</td>
<td>Recessive</td>
</tr>
<tr>
<td></td>
<td>GG</td>
<td>AG+AA</td>
</tr>
<tr>
<td>Age (years)</td>
<td>52.73±9.00</td>
<td>53.04±10.15</td>
</tr>
<tr>
<td>Sex (male/female)</td>
<td>20 (69.0)/</td>
<td>9 (31.0)/</td>
</tr>
<tr>
<td>BMI (Kg/m2)</td>
<td>24.64±4.11</td>
<td>25.89±4.54</td>
</tr>
<tr>
<td>WC (cm)</td>
<td>86.92±7.56</td>
<td>87.70±10.75</td>
</tr>
<tr>
<td>SBP (mmHg)</td>
<td>129.92±20.54</td>
<td>128.45±16.83</td>
</tr>
<tr>
<td>DBP (mmHg)</td>
<td>80.56±9.56</td>
<td>79.38±8.09</td>
</tr>
<tr>
<td>FBG (mg/dL)</td>
<td>171.12±51.64</td>
<td>192.77±72.38</td>
</tr>
<tr>
<td>HbA1c (%)</td>
<td>9.21±1.67</td>
<td>9.78±1.98</td>
</tr>
<tr>
<td>CrSr (mg/dL)</td>
<td>0.72±0.16</td>
<td>0.71±0.22</td>
</tr>
<tr>
<td>eGFR (mL/min/1.73m&lt;sup&gt;2&lt;/sup&gt;)</td>
<td>99.34±16.15</td>
<td>96.38±19.32</td>
</tr>
</tbody>
</table>

Continuous data are presented in mean±SD and categorical data as n(%)

BMI: body mass index, WC: waist circumference, SBP: systolic blood pressure, DBP: diastolic blood pressure, FBG: fasting blood glucose, CrSr: creatinine serum, eGFR: estimated glomerular filtration rate, HDL-c: high density lipoprotein cholesterol, LDL-c: low density lipoprotein cholesterol, TC: total cholesterol

Table 2. Comparison of allelic frequency of PRKAA2 rs2796498, rs9803799, and rs2746342 variants distribution in responders and non-responder of metformin therapy
<table>
<thead>
<tr>
<th>SNP ID</th>
<th>Genotype</th>
<th>Responders (n=45)</th>
<th>Non responders (n=61)</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>rs2796498</td>
<td>GG</td>
<td>22 (48.9)</td>
<td>37 (60.7)</td>
<td>0.44</td>
</tr>
<tr>
<td></td>
<td>AG</td>
<td>21 (46.7)</td>
<td>21 (34.4)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>AA</td>
<td>2 (4.4)</td>
<td>3 (4.9)</td>
<td></td>
</tr>
<tr>
<td>Alleles</td>
<td>G</td>
<td>65 (72.2)</td>
<td>95 (77.9)</td>
<td>0.35</td>
</tr>
<tr>
<td></td>
<td>A</td>
<td>25 (27.8)</td>
<td>27 (22.1)</td>
<td></td>
</tr>
<tr>
<td>HWE-p value</td>
<td></td>
<td>0.47</td>
<td></td>
<td></td>
</tr>
<tr>
<td>rs9803799</td>
<td>TT</td>
<td>37 (82.2)</td>
<td>56 (91.8)</td>
<td>0.31</td>
</tr>
<tr>
<td></td>
<td>GT</td>
<td>7 (15.6)</td>
<td>4 (6.6)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>GG</td>
<td>1 (2.2)</td>
<td>1 (1.6)</td>
<td></td>
</tr>
<tr>
<td>Alleles</td>
<td>T</td>
<td>81 (90.0)</td>
<td>116 (95.1)</td>
<td>0.15</td>
</tr>
<tr>
<td></td>
<td>G</td>
<td>9 (10.0)</td>
<td>6 (4.9)</td>
<td></td>
</tr>
<tr>
<td>HWE-p value</td>
<td></td>
<td>0.03</td>
<td></td>
<td></td>
</tr>
<tr>
<td>rs2746342</td>
<td>GG</td>
<td>12 (26.7)</td>
<td>24 (39.3)</td>
<td>0.28</td>
</tr>
<tr>
<td></td>
<td>GT</td>
<td>26 (57.8)</td>
<td>26 (42.6)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>TT</td>
<td>7 (15.5)</td>
<td>11 (18.1)</td>
<td></td>
</tr>
<tr>
<td>Alleles</td>
<td>T</td>
<td>40 (44.4)</td>
<td>48 (39.3)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>G</td>
<td>50 (55.6)</td>
<td>74 (60.7)</td>
<td>0.46</td>
</tr>
<tr>
<td>HWE-p value</td>
<td></td>
<td>0.92</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Table 3. Follow-up characteristics in drug-naive T2DM patients after receiving monotherapy metformin based on PRKAA2 genetic variation

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>rs2796498</th>
<th></th>
<th>rs9803799</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>GG</td>
<td>AG+AA</td>
<td>p-value</td>
<td>GG+AG</td>
</tr>
<tr>
<td>SBP (mmHg)</td>
<td>126.03±12.63</td>
<td>124.79±7.89</td>
<td>0.56</td>
<td>128.00±6.16</td>
</tr>
<tr>
<td>DBP (mmHg)</td>
<td>80.69±6.94</td>
<td>80.43±6.57</td>
<td>0.84</td>
<td>80.20±5.54</td>
</tr>
<tr>
<td>FBG-post (mg/dL)</td>
<td>134.66±29.85</td>
<td>139.53±28.48</td>
<td>0.40</td>
<td>155.20±36.63</td>
</tr>
<tr>
<td>HbA1c(%)</td>
<td>8.16±1.26</td>
<td>8.42±1.33</td>
<td>0.31</td>
<td>8.84±1.27</td>
</tr>
<tr>
<td>HbA1c changes (%)</td>
<td>1.03±1.18</td>
<td>1.36±1.76</td>
<td>0.25</td>
<td>1.19±1.48</td>
</tr>
<tr>
<td>HDL-c (mg/dL)</td>
<td>47.90±8.40</td>
<td>47.45±8.34</td>
<td>0.78</td>
<td>45.60±7.57</td>
</tr>
<tr>
<td>LDL-c (mg/dL)</td>
<td>104.02±24.89</td>
<td>110.43±32.71</td>
<td>0.26</td>
<td>106.16±28.70</td>
</tr>
<tr>
<td>TC (mg/dL)</td>
<td>177.25±36.12</td>
<td>183.98±34.01</td>
<td>0.33</td>
<td>196.00±30.49</td>
</tr>
<tr>
<td>Triglycerides (mg/dL)</td>
<td>140.98±61.60</td>
<td>130.45±51.07</td>
<td>0.35</td>
<td>137.74±35.77</td>
</tr>
</tbody>
</table>

Data are presented in mean±SD

BMI: body mass index, WC: waist circumference, SBP: systolic blood pressure, DBP: diastolic blood pressure, FBG: fasting blood glucose, CrSr: creatinine serum, eGFR: estimated glomerular filtration rate, HDL-c: high density lipoprotein cholesterol, LDL-c: low density lipoprotein cholesterol, TC: total cholesterol
Furthermore, multinominal logistic regression models were applied to verify the effect of \textit{PRKAA2} genetic variation on metformin efficacy (Table 4). This study failed to detect the association between \textit{PRKAA2} genetic variation with metformin efficacy, even after adjusting age, gender, and adjusting for BMI, WC, and lipid profiles. Interestingly, we found some association after adjusting for BMI, WC, lipid profiles, blood pressure, and eGFR. As shown in Table 4. AG of the rs2796498 genetic variation was statistically significant to decrease metformin efficacy (OR=0.32, 95%CI=0.12 – 0.86). Dominant model of rs2796498 (GG vs. AG+AA) showed significant association to metformin efficacy (OR=0.34, 95%CI=0.13 – 0.90).

Table 4. Association of \textit{PRKAA2} genetic variations between responders and non-responders receiving metformin therapy

\begin{tabular}{|c|c|c|c|}
\hline
Genotype & OR (95\% CI) & AOR (95\%CI)\textsuperscript{a} & AOR (95\%CI)\textsuperscript{b} \\
\hline
rs2796498 & & & \\
GG & 1.00 (reference) & & \\
AG & 0.57 (0.25 – 1.27) & 0.61 (0.27 – 1.37) & 0.32 (0.12 – 0.86)* \\
AA & 0.85 (0.13 – 5.51) & 0.90 (0.14 – 5.85) & 0.67 (0.07 – 6.50) \\
Dominant (GG vs. AG+AA) & 0.60 (0.27 – 1.30) & 0.63 (0.29 – 1.40) & 0.34 (0.13 -0.90)* \\
Recessive (GG+AG vs. AA) & 1.09 (0.17 – 6.79) & 1.12 (0.92 – 1.13) & 1.15 (0.13 – 10.46) \\
G allele & 1.00 (reference) & & \\
A allele & 0.75 (0.40 – 1.42) & 0.75 (0.40 – 1.41) & 0.55 (0.26 – 1.14) \\
rs9803799 & & & \\
TT & 1.00 (reference) & & \\
GT & 0.37 (0.10 – 1.35) & 0.37 (0.10 – 1.36) & 0.45 (0.09 – 2.16) \\
GG & 0.65 (0.04 – 10.61) & 0.95 (0.04 – 20.15) & 0.14 (0.00 – 54.13) \\
Dominant (TT vs. GT+GG) & 0.40 (0.12 – 1.33) & 0.41 (0.12 – 1.36) & 0.45 (0.11 – 1.86) \\
Recessive (TT+GT vs. GG) & 0.72 (0.04 – 11.77) & 1.05 (0.05 – 22.20) & 0.14 (0.00 – 54.62) \\
T allele & 1.00 (reference) & & \\
G allele & 0.46 (0.16 – 1.34) & 0.46 (0.16 – 1.37) & 0.52 (0.15 – 1.77) \\
rs2746342 & & & \\
GG & 1.00 (reference) & & \\
GT & 0.46 (0.19 – 1.12) & 0.52 (0.21 – 1.26) & 0.43 (0.16 – 1.19) \\
TT & 0.72 (0.22 – 2.36) & 0.83 (0.25 – 2.71) & 0.65 (0.17 – 2.50) \\
Dominant (GG vs. GT+TT) & 0.51 (0.22 – 1.21) & 0.58 (0.25 – 1.35) & 0.47 (0.18 – 1.24) \\
Recessive (GG+GT vs. TT) & 1.16 (0.41 – 3.29) & 1.24 (0.43 – 3.53) & 1.08 (0.33 – 3.55) \\
G allele & 1.00 (reference) & & \\
T allele & 0.81 (0.47 – 1.41) & 0.83 (0.48 – 1.45) & 0.78 (0.42 – 1.44) \\
\hline
\end{tabular}

\textsuperscript{a}adjusted for age and gender, \textsuperscript{b}adjusted for BMI, WC, lipid profiles, glomerular filtration rate, and blood pressure.

\section*{Discussion}

The previous study found an association between rs2796498 and rs2746342 with T2DM susceptibility [21, 22], and rs9803799 with metformin efficacy [23]. Accordingly, this current study evaluates the impact of \textit{PRKAA2} genetic variation on metformin efficacy. To the authors knowledge, this study is the first pharmacogenomic research reporting allele frequency of rs2796498, rs9803799, and rs2746342 applied metformin therapy among Indonesia drug-naïve T2DM patients.

This recent study found that T2DM patients with AA genotype of rs2796498 had significantly higher BMI than GG+AG. However, this significant mean of BMI difference was not observed in rs9803799 and rs2746342. These findings might imply that the AA genotype of rs2796498 is associated with obesity, then it should be correlated with lipid profiles. Our findings contradict the data obtained by Jones et al. [25], which indicated that rs2796498 and rs2746342 genetic variations correlated lipid profiles. Furthermore, this study found that the TT genotype of rs2746342 had higher FBG and HbA1c levels significantly than GG+GT among study subjects before receiving metformin. Conversely, Shen et al. reported that the G allele had a higher FBG level and T2DM risk [22]. It could
be caused by dissimilar ethnicity, where Shen et al. focused on Han Chinese, and this study focused on Indonesia. Nevertheless, the recessive model of rs2746342 did not influence the difference in FBG, HbA1c, and HbA1c change after metformin therapy. It could be an early indicator that rs2746342 genetic variation does not alter metformin efficacy in our population.

AMPK, as the main target of metformin, regulates the function of hepatic glucose metabolism and pancreatic b-cell [8, 11, 26]. It has been widely agreed that the phosphorylation of AMPK is induced by metformin, although the specific route is not clear yet. An animal study confirmed that 10 weeks of metformin treatment significantly escalated AMPK phosphorylation on the α2 subunit [27]. Several PRKAA2 genetic variations have been investigated to increase T2DM risk, including that’s located in the intron region [21, 22, 28]. Nevertheless, only a few studies observe that SNP is related to metformin responses [23]. A review has been mentioned that PRKAA2 genetic variation is one of the genes contributing to the metformin mechanism [20]. However, it has not yet been discovered clearly in a clinical study.

Thus, this study investigated the association of PRKAA2 genetic variation and metformin efficacy among drug-naïve T2DM patients. Our results discovered that allele and genotype frequencies of rs2796498, rs9803799, and rs2746342 between responders and non-responders were not significantly different. Interestingly, the wild type of rs2796498 (AA) and rs9803799 (GG) are detected in little number, either in the responder or non-responder group. It confirmed that those SNP has related to T2DM [21, 29], and our study participants were T2DM, indeed.

Moreover, we found a significant association between rs2796498 in AG genotype and dominant model with metformin efficacy, after adjusting for BMI, WC, lipid profiles, blood pressure, and eGFR. AG carrier in rs2796498 had 0.32 times of metformin response compared with the GG carrier. Therefore, T2DM patients with AG carriers might have a poor response to metformin therapy. Apart from these, the dominant model found that AG+AA had a worse metformin efficacy compared with GG in drug-naïve T2DM patients. However, we could not detect that the A allele influences metformin efficacy. This point emphasizes that AG carrier in rs2796498 is the predictive factors of metformin efficacy in our population. Although metformin is well-known as a better treatment for T2DM patients with obesity (ESC/EASD) [2], a different BMI in baseline could influence the impact of AA genotype in metformin efficacy.

On the other hand, we could not identify any relationship between rs9803799 and metformin efficacy. Conversely, a study in the U.S population reported that rs9803799 had significant interaction with metformin [23]. In addition, rs9803799 deviated from HWE. Therefore, findings related to rs9803799 may be reported bias association [30].

This study still has several limitations. First, the epistatic mechanism may contribute to metformin efficacy. It is possible if other genes or SNPs also affect metformin response. Second, environmental factors such as diet, physical activities, and adherence might impact decreasing HbA1c. Third, an SNP was not in agreement with HWE. Finally, the sample size should be larger to detect the function of PRKAA2 in metformin efficacy. Further replication pharmacogenomic studies are needed to confirm the PRKAA2 genetic variation on metformin efficacy.

**Conclusions**

In summary, there were no significant differences between genotype and allele frequencies of PRKAA2 genetic variation with metformin efficacy. Only AG genotype and dominant model of PRKAA2 rs2796498 associated with metformin efficacy in drug-naive T2DM patients treated with metformin as monotherapy, after adjusting for BMI, WC, blood pressure, eGFR, and lipid profiles. Nonetheless, PRKAA2 rs9803799 and rs2746342 might have no impact on metformin efficacy among Indonesian. Further study recruiting a larger sample size and engaging environment factors such as physical activities are required to confirm the role of PRKAA2 genetic variation, especially of rs2796498, rs9803799, and rs274634 on metformin efficacy among drug-naïve T2DM patients.

**Abbreviation**

- **T2DM:** type 2 diabetes mellitus
- **AMPK:** AMP-activated protein kinase
- **ACC:** acetyl-CoA carboxylase
- **PHC:** primary health care
- **FBG:** fasting blood glucose
- **BMI:** body mass index
- **WC:** waist circumference
- **SBP:** systolic blood pressure
- **DBP:** diastolic blood pressure
- **FBG:** fasting blood glucose
- **CrSr:** creatinine serum
- **eGFR:** estimated glomerular filtration rate
HDL-c: high density lipoprotein cholesterol
LDL-c: low density lipoprotein cholesterol
TC: total cholesterol

Declarations

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ETHICS APPROVAL AND CONSENT TO PARTICIPATE

The study protocol was approved by the Medical and Health Research Ethics Committee (MHREC) Faculty of Medicine, Public Health, and Nursing Universitas Gadjah Mada – Dr. Sardjito General Hospital. All measurements and data collected were done after obtaining informed consent signed by all participants.

CONSENT FOR PUBLICATION

Not applicable.

AVAILABILITY OF DATA AND MATERIALS

All data generated or analysed during this study are included in this published article (Additional file 1).

COMPETING INTERESTS

There are no conflicts of interest to declare.

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AUTHOR'S CONTRIBUTION

DV: collected patient's data in primary health care, conducted the genetic analysis, analyzed data, and wrote the manuscript. MW: designed and managed the study, wrote and corrected the manuscript. DN: interpreted the results and wrote the manuscript. All authors read and approved the final manuscript.

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References


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