

# Association between polymorphisms of *LEP*, *LEPR*, *DRD2*, *HTR2A* and *HTR2C* genes and risperidone- or clozapine-induced hyperglycemia

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**Objective:** To determine whether genetic polymorphisms related to pharmacodynamics with metabolic adverse effects, namely leptin promoter (*LEP*) rs7799039, leptin receptor rs1137101, dopamine D2 rs4436578, serotonin 5-HT2A rs6313, and serotonin 5-HT2C rs518147 and rs12836771, are associated with hyperglycemia induced by risperidone or clozapine in adult Thai patients with psychosis.

**Methods:** A total of 180 patients treated with risperidone-based ( $n=130$ ) or clozapine-based ( $n=50$ ) regimens were included in this study. Blood samples were analyzed for genotyping of the candidate genes and biochemical testing. Genotyping was performed by conducting a TaqMan real-time polymerase chain reaction-based analysis.

**Results:** The prevalence of hyperglycemia was higher in patients receiving clozapine (64.0%) than in those receiving risperidone (30.8%). Among the candidate genes, only the *LEP* rs7799039 polymorphism demonstrated a significant association with hyperglycemia ( $\chi^2=9.879$ ,  $P=0.008$ ) in patients treated with risperidone; patients with the AA genotype had the highest risk (41.1%), followed by those with AG (20.8%) and GG (0%) genotypes. Using the recessive genetic model (AA vs AG + GG), the odds ratio and 95% CI were 3.28 and 1.44–7.50, respectively. None of the genes were associated with hyperglycemia in patients treated with clozapine. A binary logistic regression revealed that the *LEP* rs7799039 polymorphism demonstrated a significant association with hyperglycemia, independent of body-mass index (BMI) in patients receiving risperidone; the odds ratio (95% CI) was 3.188 (1.399–7.262),  $P=0.006$ . By contrast, none of the pharmacodynamic genetic factors, except for BMI, were significantly associated with hyperglycemia in patients receiving clozapine.

**Conclusion:** The risk of type 2 diabetes mellitus is associated with the *LEP* rs7799039 polymorphism in Thai adults receiving risperidone but not in those receiving clozapine. Clarifying underlying mechanisms and risk of hyperglycemia provides an opportunity to prevent impaired glucose metabolism in patients receiving risperidone or clozapine.

**Keywords:** atypical antipsychotics, leptin promoter, leptin receptor, dopamine D2, serotonin 5-HT2A, serotonin 5-HT2C, diabetes mellitus

## Introduction

Second-generation antipsychotic medications can considerably ameliorate the symptoms of psychotic disorders as well as prevent relapse and treatment discontinuation.<sup>1,2</sup> However, metabolic adverse effects, particularly rapid weight gain and metabolic and endocrine abnormalities, are substantial problems for patients taking these medications.<sup>3–5</sup> Patients with schizophrenia have a high risk of chronic diseases, such as type 2 diabetes mellitus (type 2 DM) and cardiovascular disease, or other

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obesity-related complications, independent of any antipsychotic effects.<sup>6–8</sup> Consequently, patients with schizophrenia who are treated with these antipsychotic medications have a high risk of metabolic dysregulation.<sup>9,10</sup>

The prevalence of diabetes in individuals with psychotic disorders who receive treatment with antipsychotic drugs has been increasing over time; moreover, these individuals are 2–3 times more likely to develop diabetes compared with the general population.<sup>11,12</sup> Published findings from clinical trials in patients with schizophrenia who undergo treatment with clozapine and olanzapine have indicated that these medications can disrupt glucose regulation.<sup>13,14</sup> However, the results have been less conclusive for the disruption of glucose regulation for risperidone and type 2 DM; some studies have demonstrated a significantly positive association,<sup>11</sup> whereas others have reported no association.<sup>13</sup> In addition, a cross-sectional study conducted in large UK populations demonstrated that the prevalence of type 2 DM in patients with severe mental illness was higher in patients from ethnic minority groups (especially those who were Indian, Pakistani, or Bangladeshi) than in those who identified as white British.<sup>15</sup> Thus, heterogeneity in the risk of diabetes has been observed not only among individual patients but also among ethnic groups.

Mechanisms underlying antipsychotic-induced diabetes are not well understood and possibly depend on multifactorial processes.<sup>16,17</sup> Related genetic polymorphisms may explain the occurrence of hyperglycemia in some patients but not others. Different drugs interact with various neurotransmitter receptors, such as dopamine D2 and serotonin 5-HT2A and 5-HT2C; thus, a genetic susceptibility between these pharmacodynamics may contribute to glucose homeostasis.<sup>18,19</sup> Other possibilities include the role of hormones involved in glucose regulation, such as the neuroendocrine leptin.<sup>20</sup> The relationship among antipsychotic use, weight gain, metabolic disturbance, and the pharmacogenetics of leptin is still under investigation.

Several studies investigating the relationship between genetic factors and metabolic outcomes in patients treated with antipsychotics have offered inconsistent conclusions.<sup>21–24</sup> This conflict in results may be attributable to differences in study patients' characteristics, such as treatment (enrolled single/multiple types of antipsychotics) and ethnicity, suggesting the influence of genetics. Furthermore, few studies have investigated pharmacodynamic genetic factors and hyperglycemia development in patients treated with risperidone or clozapine, 2 medications commonly prescribed for both acute and long-term schizophrenia.<sup>25,26</sup> The aim of this study was to explore

whether the genetic polymorphisms of the leptin promoter (*LEP*) rs7799039, leptin receptor (*LEPR*) rs1137101, dopamine D2 (*DRD2*) rs4436578, serotonin 5-HT2A (*HTR2A*) rs6313, serotonin 5-HT2C- (*HTR2C*) rs518147, and *HTR2C* rs12836771 are associated with hyperglycemia induced by risperidone or clozapine in ethnically Thai adult patients with psychosis disorders. Identifying polymorphisms of relevant genes in patients with psychosis may help individualize treatment to prevent impaired glucose regulation.

## Materials and methods

### Study patients

This cross-sectional observational study enrolled 202 Thai patients who had received treatment for psychosis at the Somdet Chaopraya Institute of Psychiatry in central Thailand. All patients were diagnosed by a psychiatrist. Patients were informed of the specific risks and benefits of participation, and written consent was obtained from patients. The study protocol was approved by the Ethics Committee of the Faculty of Medicine, Ramathibodi Hospital, Mahidol University, and the research center of the Somdet Chaopraya Institute of Psychiatry, Bangkok, Thailand. We excluded 22 patients who were treated with antipsychotic polypharmacy ( $n=10$ ), quetiapine ( $n=4$ ), or olanzapine ( $n=8$ ). The study group consisted of 180 participants who had received either risperidone ( $n=130$ ) or clozapine ( $n=50$ ) for at least 1 month. Patient demographics and treatment history, including dosage and concomitant therapy, were extracted from medical and pharmacy records. Height, weight, and waist circumference measurements were also obtained at the time of enrollment. Exclusion criteria for this study were a known history of pervasive developmental disorders or conditions associated with convulsions, cancer, end-stage chronic kidney disease, or other serious physical conditions (ie, thyroid disorders). We excluded patients receiving lithium and sodium valproate or concomitant treatments that could potentially affect glucose and lipid metabolism.

Metabolic syndrome was defined using National Cholesterol Education Program-Third Adult Treatment Panel criteria modified for Asian populations.<sup>27</sup>

### Single-nucleotide polymorphism selection and genotyping analysis

Candidate single-nucleotide polymorphisms (SNPs) were identified from three sources. Firstly, candidate SNPs were selected based on pathophysiological pathways in

literature reviews of risperidone and clozapine-induced metabolic adverse effect such as weight gain and insulin resistance. Secondly, the minor allele frequency was greater than 0.05 in the Chinese Han in Beijing according to the International HapMap Project database (<http://hapmap.ncbi.nlm.nih.gov>). Thirdly, the significant association between the candidate gene and metabolic adverse effect in patients using antipsychotic medication has been reported in previous studies; *LEP*<sup>20,21,24,28</sup> *LEPR*<sup>20,29</sup> *DRD2*<sup>30</sup> *HTR2A*<sup>31</sup> and *HTR2C*<sup>22–24,32</sup>. In total, 6 SNPs related to pharmacodynamics genes were selected: *LEP* rs7799039 (–2548G/A), *LEPR* rs1137101 (668A/G), *DRD2* rs4436578 (Tag-SNP T/C), *HTR2A* rs6313 (102T/C), *HTR2C* rs518147 (–697G/C), and *HTR2C* rs12836771 (Tag-SNP A/G). These SNPs were genotyped by performing a TaqMan-based analysis on an Applied Biosystems®ViiA™7 real-time PCR system (ABI, Foster City, CA, USA) according to the manufacturer's protocol.

For genomic DNA extraction, blood samples were collected in EDTA tubes. DNA was isolated using the MagNA Pure automated extraction system (Roche Diagnostics), which is based on magnetic-bead technology with a lysis buffer and proteinase K. In this system, nucleic acids were bound to the surface of magnetic glass particles. Cellular debris was removed through several washing steps, and purified nucleic acids were eluted. From the 1-mL input volume of EDTA whole blood, a 200- $\mu$ L output volume of extracted genomic DNA product was obtained. The quality and purity of genomic DNA were assessed using a Nano Drop ND-1000 (NanoDrop Technologies, Wilmington, DE, USA), with dynamic ranges of 220–750 nm. Wavelengths of 260 nm and 280 nm were considered to be suitable for measuring genomic DNA and evaluating contaminated proteins in the sample, respectively. In this study, 20 ng of the purified genomic DNA template was used, and the optical density ratio at 260/280 nm was >1.7. All DNA samples were aliquoted and stored at –20 °C until analysis.

## Biochemical measurements

Blood samples were collected as serum or plasma in the fasting state. All samples were stored at 2 °C to 8 °C and analyzed within one day of collection for total cholesterol, triglyceride, low-density lipoprotein-cholesterol (LDL-C), high-density lipoprotein-cholesterol (HDL-C), glucose, insulin, adiponectin, leptin, and prolactin. Serum total cholesterol, triglyceride, LDL-C, HDL-C, and plasma glucose levels were tested using Siemens

enzymatic methods (Siemens Medical Solution Diagnostics, Tarrytown, NY, USA). The high-sensitivity C-reactive protein level was measured on the Siemens BN Prospec (Siemens Medical Solution Diagnostics) by using the immune nephelometric method, and adiponectin and leptin levels were quantified using a sandwich ELISA system (Mediagnost Gesellschaft für Forschung and Herstellung von Diagnostika GmbH, D-72770 Reutlingen, Germany). Insulin and prolactin levels were determined using a chemiluminescent immunoassay on Immulite H2975 (Siemens Medical Solution Diagnostics) and Architect ci2000 (Abbott Laboratories Abbott Park, IL, USA), respectively. Insulin resistance was estimated using the homeostasis model assessment of insulin resistance (HOMA-IR). The HOMA-IR index was calculated using the following formula:  $\text{HOMA-IR} = \text{fasting insulin } (\mu\text{U/mL}) \times \text{fasting glucose (mmol/L)} / 22.5$ .

Lipoprotein subclass patterns were analyzed using polyacrylamide tube gel electrophoresis (Quantimetrix Lipoprint™, Redondo Beach, CA, USA), which electrophoretically separates plasma lipoproteins into the following bands: very low density lipoprotein; intermediate low density lipoprotein (midband-C, midband-B, and midband-A); large-buoyant LDL (LDL1 and LDL2); small-dense LDL (LDL3 to LDL7); and HDL. Relative areas for each lipoprotein band were determined through densitometry and multiplied by the total cholesterol concentration to yield the amount of cholesterol for each band. Mean LDL particle sizes were computed.

## Statistical analyses

Continuous data for all clinical parameters and biochemical markers are reported as medians (minimum-maximum) and means (standard error of mean), and categorical variables are presented as numbers and percentages. Differences among groups were assessed using the  $\chi^2$  test, Mann–Whitney U test, or *t* test, as appropriate. A  $\chi^2$  test (Pearson chi-square or Fisher's exact test) was used to assess differences in genotype frequencies among normal fasting plasma glucose (fasting glucose <5.55 mmol/L) and increased fasting plasma glucose (fasting glucose  $\geq$ 5.55 mmol/L) groups. The recessive genetic model was used for *LEP* rs7799039: AA vs AG + GG and *LEPR* rs1137101: GG vs AG + GG and the dominant genetic model was used for *DRD2* rs4436578: CC + CT vs TT; *HTR2A* rs6313: CC + CT vs TT; *HTR2C* rs518147: CC + CG vs GG and *HTR2C* rs12836771: GG + AG vs AA. The association between

the candidate gene polymorphisms and hyperglycemia was considered by the odds ratio (OR) and the corresponding 95% confidence interval (CI). A backward, stepwise multivariable logistic regression model was used to determine whether one or more pharmacodynamic genetic factors might predict hyperglycemia when non-pharmacodynamic genetic factors (sex, age, body-mass index [BMI], smoking status, and treatment duration) were also considered. The criteria of  $P=0.05$  for a variable to enter and  $P>0.10$  for a variable to be removed.  $P<0.05$  (2 tailed) was considered statistically

significant. All analyses were performed using SPSS, version 18.0 (SPSS Inc., Chicago, IL, USA).

## Results

### Characteristics and biomarkers of the study population

The study group consisted of 180 patients (86 men and 94 women) between the ages of 18 and 77 years. Schizophrenia was diagnosed in 85.5% of patients. The demographic and biochemical test results of participants stratified by the

**Table 1** Clinical characteristics of patients treated with risperidone or clozapine

	All (n=180)	Risperidone (n=130)	Clozapine (n=50)	p-value
Clinical parameter <sup>a</sup>				
Males, n (%)	86 (47.8)	61 (46.9)	25 (50)	0.741
Age, years	42.1 (18–77)	41.0 (18–77)	44.6 (26–68)	0.073
Diagnosis, n (%)				0.016
Schizophrenia	154 (85.5)	106 (81.5)	48 (96.0)	
Other diagnosis	26 (14.5)	24 (18.5)	2 (4.0)	
Dose equivalent to olanzapine <sup>b</sup>	8.17 (0.82–25.00)	10.00 (1.25–25.00)	6.54 (0.82–13.07)	0.001
Duration of treatment, months	38.2 (1.2–227.0)	24.6 (1.2–160.7)	81.5 (3.7–227.0)	<0.001
Weight at antipsychotic initiation, kg	58.4 (32.0–118.0)	57.9 (32.0–118.0)	59.0 (37.4–95.0)	0.433
Weight at study visit, kg	62.0 (38.0–136.9)	63.0 (38.0–136.9)	60.3 (45.0–122.0)	0.670
Waist circumference, cm	87.0 (64–131)	88.5 (64–131)	87.0 (66–129)	0.767
Hip circumference, cm	96.0 (71–135)	97.0 (71–135)	94.5 (71–128)	0.094
Waist/hip ratio	0.90 (0.75–1.13)	0.89 (0.76–1.13)	0.92 (0.75–1.10)	0.028
Body mass index, kg/m <sup>2</sup>	24.9 (17.2–48.0)	25.1 (17.2–45.4)	24.2 (18.0–48.0)	0.340
Current smoker, n (%)	47 (26.1)	32 (24.6)	15 (30.0)	0.570
Alcohol consumption, n (%)	7 (3.9)	4 (3.1)	3 (6.0)	0.399
Biomarker measures <sup>c</sup>				
Fasting glucose, mmol/L	5.67 (0.11)	5.50 (0.12)	6.12 (0.27)	0.014
Insulin, $\mu$ U/mL	7.237 (0.694)	6.993 (0.773)	7.870 (1.494)	0.573
HOMA-IR	2.078 (0.302)	1.977 (0.382)	2.340 (0.447)	0.592
hsCRP, mg/L	3.166 (0.431)	3.392 (0.569)	2.677 (0.588)	0.443
Leptin, ng/mL	12.86 (0.89)	13.29 (1.05)	11.73 (1.68)	0.433
Adiponectin, $\mu$ g/mL	23.19 (1.33)	22.67 (1.44)	24.57 (3.00)	0.523
Prolactin, ng/mL	46.7 (3.4)	60.1 (4.1)	12.0 (1.5)	<0.001
Triglycerides, mmol/L	1.58 (0.10)	1.54 (0.11)	1.69 (0.21)	0.490
Total cholesterol, mmol/L	5.37 (0.09)	5.39 (0.10)	5.31 (0.19)	0.703
HDL-C, mmol/L	1.47 (0.03)	1.49 (0.04)	1.41 (0.06)	0.279
LDL-C, mmol/L	3.37 (0.08)	3.39 (0.09)	3.31 (0.15)	0.654
Small-dense LDL, mmol/L	0.33 (0.03)	0.34 (0.04)	0.33 (0.08)	0.920
Mean LDL particle size, nm	26.66 (0.46)	26.66 (0.52)	26.66 (0.94)	0.999

**Notes:** <sup>a</sup>Data except for number (%) are the median (minimum-maximum). <sup>b</sup>The doses equivalent to olanzapine are estimated according to the publication by Leucht et al<sup>33</sup> risperidone 0.4 mg/d or clozapine 30.6 mg/d is equal to olanzapine 1 mg/d. <sup>c</sup>Data are given as mean (standard error of mean). All biochemical markers are expressed in Système international units; conversions to conventional units are as follows: fasting glucose (mg/dL), multiply by 18.02; triglycerides (mg/dL), multiply by 88.5; cholesterol (mg/dL), multiply by 38.6.

**Abbreviations:** HOMA-IR, homeostasis Model Assessment of Insulin Resistance; hsCRP, high sensitivity C-reactive protein; HDL-C, high density lipoprotein-cholesterol; LDL-C, low density lipoprotein-cholesterol; LDL, low density lipoprotein.

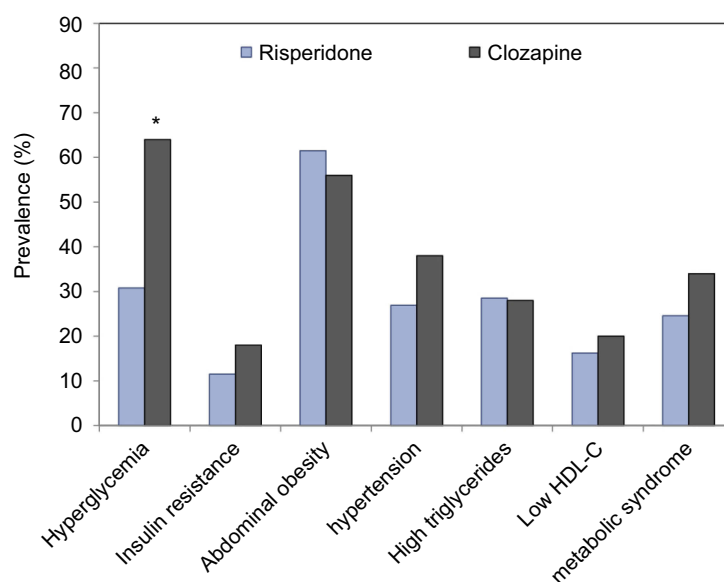
medication prescribed are summarized in Table 1. Overall, 130 (72.2%) and 50 (27.8%) patients were receiving risperidone and clozapine treatment, respectively. For patients in our study, the median dose equivalent to olanzapine<sup>33</sup> for risperidone treatment (10.00 mg/d; interquartile range: 5.00–15.00 mg/d) was significantly higher than that for clozapine treatment (6.54 mg/d; interquartile range: 3.27–9.80 mg/d). By contrast, the duration of clozapine treatment was significantly longer than that of risperidone treatment; the median treatment durations for clozapine and risperidone were 81.5 (interquartile range: 25.0–110.8 months) and 24.6 months (interquartile range: 10.8–51.2 months), respectively. Patients treated with each antipsychotic medication were similar with respect to sex, age, weight, BMI, and prevalence of cigarette smoking and alcohol consumption. Although the median waist circumference and hip circumference did not differ between the 2 treatment groups, the waist to hip ratio was slightly higher in patients treated with clozapine than in those treated with risperidone. The results obtained for metabolic biomarkers were similar for participants in the 2 groups, with the exception of a higher mean fasting plasma glucose concentration and a lower mean prolactin concentration in patients receiving clozapine.

As indicated in Figure 1, the prevalence of hyperglycemia was higher in patients receiving clozapine [32/50 (64.0%)] than in those receiving risperidone [40/130 (30.8%)] ( $\chi^2=16.615$ ,  $P<0.001$ ). By contrast, the prevalence of insulin resistance (HOMA-IR >3.0) in patients receiving

clozapine [9/50 (18.0%)] was not significantly different from that in patients receiving risperidone [15/130 (11.5%)] ( $\chi^2=1.305$ ,  $P=0.327$ ). No difference was noted between the 2 treatment groups in terms of other clinical features of metabolic syndrome (all  $P>0.200$ ).

## Associations of pharmacodynamic gene polymorphisms and hyperglycemia

Genotype distributions in patients with normal and increased fasting plasma glucose levels are presented in Table 2. Genotype frequencies were not significantly different between patients receiving risperidone or clozapine (all  $P>0.200$ ). Among candidate genes, only the *LEP* rs7799039 polymorphism demonstrated a significant association with hyperglycemia (Pearson chi-square =9.879,  $P=0.008$ ) in risperidone-treated patients; patients with the AA genotype had the highest risk [30/73 (41.1%)], followed by those with AG [10/48 (20.8%)] and GG [0/9 (0%)] genotypes. Using the recessive genetic model (AA vs AG + GG), the odds ratio (OR) and 95% CI were 3.28 and 1.44–7.50, respectively. Although the *LEPR* rs1137101 polymorphism was not significantly associated with hyperglycemia (Fisher's exact =2.063,  $P=0.427$ ), the GG genotype exhibited the highest risk of hyperglycemia [30/91 (33.0%)], followed by AG [10/35 (28.6%)] and AA [0/4 (0%)] genotypes. None of the study genes were associated with hyperglycemia in clozapine-treated patients.



**Figure 1** The prevalence of metabolic abnormality in patients receiving risperidone or clozapine. Metabolic syndrome and its major components are defined using the NCEP ATP III criteria modified for Asian population.<sup>27</sup> \* $p$ -value<0.05.

**Table 2** Genotype frequency of *LEP*, *LEPR*, *DRD2*, *HTR2A*, and *HTR2C* polymorphisms among patients with and without impaired fasting glucose

	Risperidone				Clozapine			
	NFG <sup>a</sup> (n=90)	IFG <sup>b</sup> (n=40)	p-value <sup>c</sup>	OR (95% CI) <sup>d</sup>	NFG <sup>a</sup> (n=18)	IFG <sup>b</sup> (n=32)	p-value <sup>c</sup>	OR (95% CI) <sup>d</sup>
<i>LEP</i> rs7799039 (G>A)	9 (100.0)	0 (0.0)	0.008	3.28 (1.44–7.50)	2 (40.0)	3 (60.0)	0.913	0.78 (0.24–2.48)
	38 (79.2)	10 (20.8)			7 (31.8)	15 (68.2)		
	43 (58.9)	30 (41.1)			9 (39.1)	14 (60.9)		
<i>LEPR</i> rs1137101 (A>G)	4 (100.0)	0 (0.0)	0.427	1.92 (0.84–4.36)	0 (0.0)	1 (100.0)	>0.999	0.73 (0.19–2.82)
	25 (71.4)	10 (28.6)			4 (33.3)	8 (66.7)		
	61 (67.0)	30 (33.0)			14 (37.8)	23 (62.2)		
<i>DRD2</i> rs4436578 (T>C)	45 (72.6)	17 (27.4)	0.116	1.35 (0.64–2.87)	8 (44.4)	10 (55.6)	0.220	1.76 (0.53–5.80)
	37 (72.5)	14 (27.5)			7 (25.9)	20 (74.1)		
	8 (47.1)	9 (52.9)			3 (60.0)	2 (40.0)		
<i>HTR2A</i> rs6313 (T>C)	36 (67.9)	17 (32.1)	0.964	0.90 (0.42–1.92)	9 (37.5)	15 (62.5)	>0.999	1.13 (0.36–3.60)
	43 (69.4)	19 (30.6)			7 (33.3)	14 (66.7)		
	11 (73.3)	4 (26.7)			2 (40.0)	3 (60.0)		
<i>HTR2C</i> rs518147 (G>C)	70 (70.0)	30 (30.0)	0.269	1.17 (0.49–2.79)	13 (39.4)	20 (60.6)	0.565	1.56 (0.45–5.47)
	14 (77.8)	4 (22.2)			4 (40.0)	6 (60.0)		
	6 (50.0)	6 (50.0)			1 (14.3)	6 (85.7)		
<i>HTR2C</i> rs12836771 (A>G)	73 (70.9)	30 (29.1)	0.332	1.43 (0.59–3.48)	14 (37.8)	23 (62.2)	0.889	1.37 (0.35–5.30)
	11 (73.3)	4 (26.7)			3 (37.5)	5 (62.5)		
	6 (50.0)	6 (50.0)			1 (20.0)	4 (80.0)		

Notes: <sup>a</sup>NFG: normal fasting plasma glucose. <sup>b</sup>IFG: increased fasting plasma glucose. <sup>c</sup> $\chi^2$  test (Pearson chi-square or Fisher's exact) for the genotype frequencies between NFG and IFG groups <sup>d</sup>Odds ratio (95% confidence interval): Recessive genetic model; *LEP* rs7799039: AA vs AG + GG; *LEPR* rs1137101: GG vs AG + GG; Dominant genetic model; *DRD2* rs4436578: CC + CT vs TT; *HTR2A* rs6313: CC + CT vs TT; *HTR2C* rs518147: CC + CG vs GG; *HTR2C* rs12836771: GG + AG vs AA.

To assess the association between pharmacodynamic genetic factors and nonpharmacodynamic genetic factors with hyperglycemia in patients treated with risperidone or clozapine, we evaluated their contributions by performing a binary logistic regression, as shown in Table 3. In Model 1, all candidate genes and the clinical factors of sex, age, BMI, smoking status, and treatment duration were considered as independent variables. Individual variables that did not exhibit a statistically significant correlation ( $P>0.05$ ) with hyperglycemia were removed using a backward stepwise logistic regression; the remaining variables are presented with Model 2. *LEP* rs7799039 was the only polymorphism that demonstrated a significant association with hyperglycemia, independent of BMI in patients on risperidone (OR, 3.188; 95% CI, 1.399–7.262;  $P=0.006$ ). In patients treated with clozapine, none of the candidate genes demonstrated an association with hyperglycemia. BMI was significantly associated with hyperglycemia; the OR (95% CI) was 1.6233 (1.170–2.252).

## Discussions

Thai patients treated with clozapine demonstrated a higher rate of elevated fasting glucose (up to 64.0%), 2 times greater than that of patients treated with risperidone (30.8%). The pharmacoepidemiological–pharmacodynamic study performed using

the VigiBase database revealed that among patients with schizophrenia who had diabetes, 53.4%, 30.6%, and 7.8% had received clozapine, olanzapine, and risperidone treatment, respectively.<sup>34</sup> We noted that the rate of insulin resistance induced by clozapine and risperidone was not significantly different, suggesting that the diabetogenic effects of these drugs may operate under different mechanisms. The pharmacodynamic characteristics of each antipsychotic medication may provide further insights into mechanisms underlying the pathophysiology of glucose disturbances.

Most published studies on genetic polymorphisms have reported an association with metabolic syndrome, especially weight gain, in patients treated with antipsychotic agents.<sup>20–24,28–32,35–38</sup> Because *LEP*, *HTR2A*, and *HTR2C* are involved in energy expenditure, polymorphisms in genes coding for these receptors might affect metabolic homeostasis in patients receiving long-term treatment with antipsychotic medications.<sup>20,29–32,37</sup> However, less is known regarding their effect on type 2 DM in patients treated with antipsychotic medications. Our results indicated that *LEP* rs7799039 had a significant association with hyperglycemia in patients treated with risperidone. However, the association was independent of BMI, suggesting that risperidone-induced glycaemic dysregulation occurs irrespective of changes in

**Table 3** Logistic regression analysis between the pharmacodynamic genetic and clinical risk factors with impaired fasting glucose

Dependent Variables	Independent Variables	Logistic Regression <sup>a</sup>			
		Risperidone		Clozapine	
		Odds Ratio (95% CI)	p-value	Odds Ratio (95% CI)	p-value
Model 1 <sup>b</sup>	<i>LEP</i> rs7799039	4.168 (1.584–10.966)	0.004	4.832 (0.638–36.615)	0.127
	<i>LEPR</i> rs1137101	1.083 (0.409–2.869)	0.872	0.833 (0.065–10.732)	0.888
	<i>DRD2</i> rs4436578	1.618 (0.834–3.139)	0.155	28.618 (1.530–535.40)	0.025
	<i>HTR2A</i> rs6313	0.621 (0.286–1.347)	0.228	0.120 (0.015–0.963)	0.046
	<i>HTR2C</i> rs518147	0.523 (0.118–2.326)	0.395	12.012 (0.891–162.0)	0.061
	<i>HTR2C</i> rs12836771	1.736 (0.388–7.757)	0.470	0.123 (0.008–1.966)	0.138
	Sex	2.781 (0.939–8.237)	0.065	3.263 (0.260–40.953)	0.360
	Age	1.032 (0.989–1.077)	0.144	1.149 (0.996–1.327)	0.057
	BMI	0.951 (0.867–1.042)	0.281	2.169 (1.190–3.953)	0.012
	Smoking	0.412 (0.125–1.361)	0.146	0.016 (0.000–0.877)	0.043
	Duration of treatment	1.012 (0.994–1.029)	0.187	0.982 (0.954–1.011)	0.217
	Constant	-	0.038	-	0.021
Model 2 <sup>c</sup>	<i>LEP</i> : rs7799039	3.188 (1.399–7.262)	0.006	-	-
	BMI	-	-	1.623 (1.170–2.252)	0.004
	Smoking	-	-	0.052 (0.004–0.656)	0.002
	Constant	-	0.001	-	0.082

**Notes:** <sup>a</sup>Logistic Regression parameters; Odds ratio (95% CI): odds ratio (95% confidential interval). <sup>b</sup>Data are from logistic regression analyses: enter method; adjusted for sex, age, BMI, smoking status and duration of treatment. <sup>c</sup>Data are from logistic regression analyses: backward likelihood ratio method; adjusted for sex, age, BMI, smoking status and duration of treatment.

weight. Thus, risperidone may induce hyperglycemia through direct or indirect effects on leptin action and the leptin gene, rather than as a result of secondary weight gain. By contrast, none of the study gene polymorphisms, except for BMI, were significantly associated with hyperglycemia in patients receiving clozapine. These findings suggest that risperidone and clozapine possibly affect glucose homeostasis through different mechanisms, through a genetic polymorphism for risperidone-induced hyperglycemia and through weight gain for glucose intolerance induced by clozapine therapy.

Considering the distribution of genetic variants of *LEP* rs7799039, namely wild-type (AA), heterozygous (AG), and homozygous (GG), we discovered that on average, patients with the AA genotype had a 2-fold higher probability of developing glycemic abnormalities than did those with the AG genotype (as indicated by increased fasting glucose; 41.1% vs 20.8%). None of the patients with the GG genotype were noted to have risperidone-induced hyperglycemia. The carriers of the AA genotype demonstrated an increased risk of type 2 DM after adjustment for sex, age, BMI, smoking status, and treatment duration (OR, 3.188; 95% CI, 1.3990–7.262). Therefore, the *LEP* AA genotype may be a key genetic predictor of hyperglycemia in patients treated with risperidone, whereas the *LEP* GG genotype may not or may even be protective.

The diabetogenic effect of risperidone may be explained, at least in part, by its action on leptin and insulin resistances as well as the functionality of the *LEP* gene. Leptin acts by directly regulating pancreatic  $\beta$  cells and insulin-sensitive tissues, independent of its effects on adiposity.<sup>39</sup> In our previous study on children and adolescents with autistic spectrum disorders, we demonstrated that risperidone therapy directly and adversely affects glucose homeostasis by inhibiting the actions of leptin and insulin.<sup>40,41</sup> Studies on mice conducted by Cheng et al revealed that palmitic acid and arachidonic acid can inhibit leptin signaling in the paraventricular nucleus of the hypothalamus, attenuating the central leptin regulation of glucose homeostasis.<sup>42,43</sup> According to a study on metabolic profiling in patients with schizophrenia, treatment with risperidone may contribute to increased palmitic acid and arachidonic acid levels.<sup>44</sup> Thus, the occurrence of these metabolic disturbances after risperidone treatment is probably the cause of leptin resistance. In addition, risperidone may induce the leptin and insulin resistance in peripheral tissues, particularly skeletal muscle, through

suppression of cytokine signaling (SOCS), a critical regulator of leptin and insulin signaling pathways.<sup>45</sup> Increases in SOCS3 and SOCS6 mRNA expression levels result in the reduction of both insulin-induced protein kinase B activation and leptin-stimulated signal transducer and activator of transcription 3 phosphorylation, which also mediates insulin and leptin resistance.

The *LEP* rs7799039 (–2548G/A) is located in the promoter region of the *LEP* gene on chromosome 7q31.3.<sup>46</sup> The gene polymorphism *LEP* rs7799039 has a strong influence on leptin gene transcription, expression, and adipose tissue secretion. Numerous studies have focused on the classic influence of leptin on obesity and obesity-related diseases, including antipsychotic-induced weight gain.<sup>28,47–50</sup> Although a mutation in *LEP* rs7799039 has received recognition as a potential contributor to antipsychotic-associated weight gain, findings have yielded somewhat conflicting results.<sup>28,47,48</sup> Lee and Bishop demonstrated that both G and A alleles have significant associations with weight gain that depend on the population and antipsychotic used.<sup>28</sup> Their results indicated that a multifactorial network of pathways influence metabolic outcomes in patients treated with antipsychotic medications.

A meta-analysis study conducted by Shen et al indicated that genetic differences in the *LEP* rs7799039 polymorphism were discernible among ethnic populations.<sup>47</sup> The frequencies of G allele were high in European populations, but low in Asian populations, with AA predominating. For the Thai population in our study, genetic frequencies were approximately 56% for AA, 37% for AG, and 7% for GG genotypes. Shen et al suggested that differences in ethnicity appear to correspond to inconsistencies in the relationship between the *LEP* rs7799039 polymorphism and antipsychotic-induced weight gain.<sup>47</sup> The A allele was proposed as a genetic risk factor for weight gain in the Asian population, whereas it is considered to be a genetic protective factor in the European population. Consequently, the *LEP* rs7799039 polymorphism might increase the risk of type 2 DM in patients treated with risperidone, with effects that vary by ethnicity. As mentioned above, both the G and A alleles may differ in the function of the *LEP* gene. In regards to the gene frequencies, the recessive genetic model (AA vs AG +GG) dominated in our population, therefore the A allele may influence decreasing leptin transcription, expression and secretion. However, in our results, the mean of leptin level in the AA genotype carriers (13.54±12.97 ng/mL)



were not significantly lower than that in the AG (11.98 ±9.9 ng/mL) or GG (18.28±13.09 ng/mL) genotype carriers. Hoffstedt et al studied in Swedish, they reported the *LEP* rs7799093 polymorphism appears to influence leptin transcription, expression and secretion such that AA genotype carriers have twice the rate of leptin secretion and 60% more leptin mRNA transcription than G allele carriers.<sup>51</sup> Therefore, the dominant genetic model (AA + AG vs GG) usually dominates in European population, with the G allele influencing the function of the *LEP* gene and perhaps increasing the risk of type 2 DM in risperidone-treated patients.

Our results demonstrated that the *LEP* rs7799039 polymorphism was not associated with hyperglycemia in clozapine treated patients. The underlying mechanisms to explain less association of clozapine with the polymorphisms than risperidone is not clear. However, we believe that it may be related to variety of other genes and various environmental factors. There were many reports describing both leptin and serotonin 5-HT<sub>2C</sub> that have strong affinity for and antagonize the serotonin 5-HT<sub>2C</sub> receptors.<sup>52,53</sup> These neuropeptides act through the hypothalamic satiety mechanisms. Leptin has been shown to stimulate central serotonin turnover. Central leptin-induced anorexia is in part mediated by the 5-HT<sub>2C</sub> receptor as well. In addition, in antipsychotics treatment, the patients with the polymorphism on levels of 5-HT<sub>2C</sub> receptor expression tended to have higher baseline leptin levels and subsequent weight gain and metabolic syndrome.<sup>54</sup> Thus, it is plausible that the multifactorial network of pathways has an interaction between both neurotransmitters.<sup>55</sup> Because of linking *HTR2C* genotype with *LEP* genotype, the *HTR2C* polymorphism owing to inherent differences in gene expression and leptin signaling which may attenuate a role of leptin involved in the anorexic effect and the glucose regulatory during drug exposure.<sup>55</sup> Due to clozapine have a higher affinity to 5-HT<sub>2C</sub> receptor compared to risperidone, the functionality of polymorphism on levels of 5-HT<sub>2C</sub> receptor expression may somehow more contribute to leptin dysregulation. Therefore, the relationship between leptin levels and levels of 5-HT<sub>2C</sub> receptor expression may explain why *LEP* rs7799039 polymorphism was not associated with hyperglycemia in clozapine treated patients. Future pharmacogenetic investigation will identify mechanistically and therapeutically informative of interactions or differential associations when *HTR2C* and *LEP* genetic variants are analyzed together.

A strong association between the *LEPR* rs1137101 polymorphism and obesity has been reported.<sup>20</sup> In the present study, we did not identify an association between *LEPR* polymorphisms and the risk of diabetes. Our findings agree with those of a study published by Kuo and colleagues who demonstrated that the *LEP* gene, but not *LEPR* gene, was significantly associated with the insulin level and HOMA-IR in Taiwanese patients with schizophrenia.<sup>29</sup> However, all patients who carried the wild-type gene, approximately 3% of our study population, demonstrated normal glucose homeostasis. Thus, the presence of the homozygous A allele may be associated with a low risk of glucose dysregulation in patients treated with risperidone.

Many studies have reported that serotonin 5-HT<sub>2C</sub> receptors, which are known to be associated with antipsychotic-induced weight gain due to a high degree of occupancy with antipsychotic medications (particularly clozapine), have a strong influence on the development of glucose intolerance and diabetes.<sup>32,33,56</sup> Hypothalamic 5-HT<sub>2C</sub> receptors can mediate glucose homeostasis through the cholinergic neurons of the sympathetic nervous system, which subsequently activate systems controlling plasma glucose in adipose, muscle, and liver tissues.<sup>57</sup> In the current study, we did not identify statistically significant associations between *HTR2C* rs518147 or *HTR2C* rs12836771 (Tag-SNP of rs3813929) and hyperglycemia development in patients treated with clozapine; this finding is consistent with those of 2 studies on Caucasian and Korean patients.<sup>36,58</sup> However, BMI demonstrated a strong association with hyperglycemia development in patients treated with clozapine. This may confirm that obesity-related insulin resistance may be the cause of the diabetogenic effect seen with clozapine.<sup>11,20,59</sup>

This study has some limitations. First, this study adopted a cross-sectional design, which may not be capable of adequately confirming associations between risperidone or clozapine therapy and changes in glucose metabolism. Second, our sample size was relatively small, particularly for patients treated with clozapine. The small sample for clozapine treatment was reflected the clinical practice of psychotic patients in our population. A relatively small sample size increases the risk of a type I error with false-positive findings. Our results require validation with a larger independent sample of patients, which can provide sufficient statistical power to infer the involvement of the *LEP* gene in risperidone-induced hyperglycemia. Third, all patients in the present study were ethnically Thai. The genotype frequency of the *LEP*

rs7799039 polymorphism differ noticeably among ethnic populations. Thus, the results of this study may not be applicable to people of other ethnicities until they receive validation from a prospective study involving other non-Asian patients. Fourth, age, sex, BMI, smoking status and duration of treatment are known to be risk factors associated with developing hyperglycemia; although these factors were accounted for and adjusted in our analysis, other factors were not accounted for, despite being known to be risk factors in developing hyperglycemia in patients with schizophrenia such as the duration of untreated psychosis or duration of illness, the duration and type of previous medication, dietary intake, level of education, physical activity and family history.

In conclusion, in adult Thai patients with psychosis, we identified substantially different mechanisms underlying the effects of risperidone and clozapine on hyperglycemia. The high interindividual variability in the risk of hyperglycemia suggests that genetics may play a crucial role in an individual's susceptibility to hyperglycemia, making it a target for future research in pharmacogenetics. The polymorphism *LEP* rs7799039 was associated with risperidone-induced hyperglycemia. The *LEP* AA genotype may be a key genetic predictor of hyperglycemia, whereas the *LEP* GG genotype may be a genetic protective factor. Testing for *LEP* rs7799039 in patients with psychosis may help to individualize treatment and prevent impaired glucose metabolism in patients receiving risperidone. However, we did not observe an association between clozapine and polymorphisms assessed in this study. Clozapine-induced weight gain appeared to have a primary effect on the development of glucose intolerance and diabetes. Understanding underlying mechanisms and risks may lead to effective interventions for preventing impaired glucose metabolism in patients receiving risperidone or clozapine.

## Ethics approval and consent to participate

The study protocol was reviewed and approved by the ethics committee of Ramathibodi Hospital (Approval reference number ID MURA2015/23). All methods were carried out in accordance with the Declaration of Helsinki.

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## Author contributions

All the authors participated in the interpretation and the review of the data. PS, AP and WU designed the study. PS, AP, WU, NJ, SP and CNN conducted the data retrieval and analyzed the data. PS, AP, SV and MHK wrote the manuscript. PS, AP, CS and MHK gave constructive suggestions during the preparation of the manuscript. All the authors also participated in the revision of the manuscript and read and approved the final manuscript. All authors agree to be accountable for all aspects of the work.

## Disclosure

The abstract of this paper was presented at the 70th AACC Annual Scientific Meeting as a poster presentation with interim findings. The poster's abstract was published in "Abstract Guide" in poster sessions: Endocrinology/Hormones. All authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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