Measurement of the levels of leptin, BDNF associated with polymorphisms LEP G2548A, LEPR Gln223Arg and BDNF Val66Met in Thai with metabolic syndrome

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Abstract

Background: Metabolic syndrome is a cluster of metabolic risk factors including dyslipidemia, impaired glucose tolerance, hypertension and central obesity. BDNF (Brain-derived neurotrophic factor) and leptin have been implied in the energy homeostasis. The purposes of this study were to examine concentrations of leptin, BDNF and biochemical parameters in metabolic-syndrome subjects and healthy controls, and also to search for associations of leptin gene (LEP) G2548A, leptin receptor gene (LEPR) Gln223Arg, and BDNF gene (BDNF) Val66Met polymorphisms with leptin levels, BDNF levels and metabolic syndrome among Thais.

Methods: The case-controlled design was performed using 322 Thai volunteers (160 metabolic-syndrome subjects; 162 controls) during the health screening program. Metabolic syndrome was assessed by using the modified National Cholesterol Education Program, Adult Treatment Panel III criteria. The levels of leptin, BDNF, insulin, glucose and lipids were measured in samples. Genotyping of LEP G2548A, LEPR Gln223Arg and BDNF Val66Met was carried out using polymerase chain reaction-restriction fragment length polymorphism technique.

Results: Serum leptin levels were significantly higher in the metabolic-syndrome group than the control group (p < 0.01), but the BDNF difference between them was not significant. Significant associations of LEPR Gln223Arg polymorphism were found with leptin and glucose levels (p < 0.05), after adjusting for potential covariates. This LEPR polymorphism in the metabolic-syndrome group was also significantly more frequent than in the control group (p < 0.05). However, other gene polymorphisms, LEP G2548A and BDNF Val66Met, showed no significant relationship with leptin levels, BDNF levels or metabolic syndrome.

Conclusion: These findings suggest leptin levels are linked with metabolic syndrome. LEPR Gln223Arg polymorphism impacted leptin concentrations, and this gene polymorphism may influence susceptibility to metabolic syndrome among Thais.

Keywords: Leptin, BDNF, Gene polymorphisms, Metabolic syndrome

Introduction

Metabolic syndrome (MS), which involves abdominal obesity, dyslipidemia, hypertension, and glucose tolerance represents one of the world’s most challenging health problems, and is an important risk factor for type 2 diabetes and cardiovascular diseases [1]. Leptin, an adipocytokine, is a key metabolic regulator and acts to reduce food intake, increase energy expenditure, and modulate immune and inflammatory responses by binding and activating the leptin receptor [2]. Serum leptin is proportional to body adiposity [3]. There are several studies indicating links of leptin gene (LEP) and leptin receptor gene (LEPR) variations with obesity [4], insulin resistance [5], and type 2 diabetes [6]. The gene polymorphisms of LEP G2548A (rs7799039: guanine > adenine) and LEPR Gln223Arg (rs1137101: glutamine > arginine)
are studied widely because \( \text{LEP} \, G2548A \) has been associated with leptin production and secretion [7], and \( \text{LEPR} \, \text{Gln223Arg} \) has been associated with impaired signaling capacity of the leptin receptor [8]. However, evidence for an association of \( \text{LEP} \, G2548A \) and \( \text{LEPR} \, \text{Gln223Arg} \) with leptin concentrations and obesity and diabetes [12,13,14]. Previous studies have shown links between molecules that mediate the function in the control of energy balance. The associations of BDNF and leptin with functions involved in energy homeostasis have been studied [12]. Previous studies have shown links between BDNF concentrations and obesity and diabetes [13,14].

The most common BDNF gene (\( \text{BDNF} \)) polymorphism, Val66Met (rs6265: valine > methionine), affects intracellular trafficking of pro-BDNF, and also decreases secretion and production of the mature BDNF with the Met allele [15]. However, previous studies have shown inconsistent results in relation to BDNF concentrations in MS subjects [16,17], and information on \( \text{BDNF} \, \text{Val66Met} \) gene polymorphism and MS is scarce. Therefore, the aims of this study were to examine concentrations of leptin, BDNF and biochemical parameters in MS subjects and healthy controls, and to search for links of \( \text{LEP} \, G2548A \), \( \text{LEPR} \, \text{Gln223Arg} \) and \( \text{BDNF} \, \text{Val66Met} \) gene polymorphisms with leptin levels, BDNF levels and MS among Thais.

Brain-derived neurotrophic factor (BDNF) is a member of the neurotrophic factor family and is one of the molecules that mediate the function in the control of energy balance. The associations of BDNF and leptin with functions involved in energy homeostasis have been studied [12]. Previous studies have shown links between BDNF concentrations and obesity and diabetes [13,14].

Methods
Following approval by the Ethics Committee of Rangsit University (RSEC No.016/53), Thailand and in accordance with the Declaration of Helsinki, this case-control study enrolled 322 Thai volunteers aged between 24 to 64 years and living in urban and suburban residential areas of Bangkok, Thailand. Among them, 162 healthy controls (86 male, 76 female) and 160 MS subjects (73 male, 87 female) were chosen during the health screening program among check-up subjects in March 2011-February 2012 and the statistical power in our sample size calculation was 80% at alpha =0.05. A physical examination and medical history check on all subjects was performed. Subjects with a history of liver, kidney and cardiovascular diseases were excluded from the study. Metabolic syndrome was defined using the modified National Cholesterol Education Program/Adult Treatment Panel III (NCEP/ATP III) criteria [18]. The new cut-off for waist circumference in the Asia-Pacific Region, instead of the original cut-off in the ATP III criteria, was used. Adoption of the new cut-off point for fasting plasma glucose criterion has already been reported (> 100 mg/dl). The modified NCEP/ATP III definition required at least three of the following: raised triglyceride (TG) levels: > 150 mg/dl, reduced high-density lipoprotein cholesterol (HDL-C): < 40 mg/dl in men, and < 50 mg/dl in women, raised fasting plasma glucose: > 100 mg/dl, raised waist circumference (WC): > 90 cm in Asian men and > 80 cm in Asian women or raised blood pressure: systolic blood pressure > 130 mmHg or diastolic blood pressure > 85.

Anthropometric measurements
The body weight of each subject was measured using a carefully calibrated beam balance (Detecto®, Cardinal Detecto Scale Manufacturing, MO). Height was measured using a vertical measuring rod. Body mass index (BMI) was conventionally expressed as weight (kg)/height (m²). Waist and hip circumferences were measured. Waist and hip circumferences were calculated for waist and hip circumference ratio. Furthermore, this study had also determined triceps skin-fold thickness (TSF) and arm circumference (AC) because TSF was suggested as the simple predictor of body density and hence percentage total body fat [19] and AC measurement could reflect adult nutritional status as defined by BMI [20]. The upper arm length midpoint mark was used to measure TSF and AC. The TSF was measured by using the Holtain Skinfold Caliper. Blood pressure (BP) was measured by a nurse after 5 to 10 minutes’ rest in the sitting position.

Laboratory measurements
10 ml of venous blood was taken from subjects in the morning after overnight fast. Glucose, TG, HDL-C levels were measured using enzymatic methods by DADE Dimension AR®. Leptin and insulin levels were measured using a radioimmunoassay kit from Linco Research (St Louis, USA). Serum BDNF levels were determined using an ELISA protocol, according to the manufacturer’s instructions (DBD00; R & D Systems, Europe).

Polymerase chain reaction-restriction fragment length polymorphism (PCR-RFLP) technique
DNA was extracted from EDTA-treated whole blood by FlexiGene DNA kit (Qiagen, Hilden, Germany). Genotyping of \( \text{BDNF} \, \text{Val66Met} \), \( \text{LEP} \, G2548A \) and \( \text{LEPR} \, \text{Gln223Arg} \) was carried out using PCR-RFLP assay. DNA fragments were amplified by PCR (PE Applied Biosystems). For analysis of the \( \text{BDNF} \, \text{Val66Met} \) polymorphism, the following primers were used:

Forward primer – 5′-ATCCGAGGACAAAGGTGGG-3′
Reverse primer – 5′-CCTATGGACATGGTGCAG-3′

A 50 µl PCR reaction was performed according to the protocol described by Matsushita et al. [21]. For analysis
of the LEP G2548A polymorphism, the following primers were used:
Forward primer – 5′-TTTCTGTAATTTTCCCGTG AG-3′
Reverse primer – 5′-AAAGCAAAGACAGGCATA AAAA-3′
A 50 μl PCR reaction was conducted, according to the protocol described by Boumaiza et al. [10]. For analysis of the LEPR Gln223Arg polymorphism, the following primers were used:
Forward primer – 5′-ACCCTTTAAGCTGGGTTCAGAAATAG-3′
Reverse primer – 5′-AGCTAGCAAATATTTTTGTA AGCAATT-3′
A 50 μl PCR reaction was conducted, according to the protocol described by Duarte et al. [4]. PCR products (304 bp for BDNF, 242 bp for LEP and 421 bp for LEPR) were detected on 2% agarose gel containing ethidium bromide. Aliquots of the PCR products were digested with 8 U PmII, HhaI and MspI restriction enzymes for the BDNF Val66Met, the LEP G2548A, and the LEPR Gln223Arg polymorphisms, respectively. Alleles were visualized as fragments by electrophoresis through an ethidium bromide-stained 2.5% agarose gel. Distribution patterns of the BDNF Val66Met, the LEP G2548A, and the LEPR Gln223Arg polymorphisms were shown in Figures 1, 2, 3.

**Statistical analysis**
Statistical analysis was performed using SPSS for Windows version 11.5 (SPSS, Chicago, IL). Median, range and 95% confidence intervals (CI) were calculated. The difference between the two groups was compared by Mann-Whitney U/Wilcoxon Rank Sum W test. The differences in genotypic and allelic frequencies of the two groups were assessed by Chi-square test. The Minitab statistical computer program was used to calculate the odds ratio (OR). Spearman rank was used to calculate correlations among the variables. To assess links between MS...
as a dependent variable and other potential factors, logistic regression was applied. A p-value < 0.05 was considered statistically significant. Goodness-of-fit of the logistic regression models was established by Hosmer-Lemeshow test.

Results
Anthropometric and biochemical data of the MS and control groups are shown in Table 1. Serum leptin levels were significantly higher in the MS group than the control group (p < 0.01), while lower HDL-C levels were observed in MS group as compared to the control group (p < 0.01). Meanwhile, there were significant differences between the MS and control groups in glucose, insulin, TG, systolic BP, diastolic BP, BMI, waist/hip ratio, WC, AC and TSF (p < 0.01). The BDNF difference between the MS and control groups was not significant. Spearman's Rank correlation test results are shown in Table 2. Leptin level was positively correlated with age, insulin, TG, systolic BP, diastolic BP, BMI, waist/hip ratio, WC, AC and TSF, but negatively correlated with HDL-C level (p < 0.05). Moreover, BDNF level correlated significantly with age, AC and TSF, as shown in Table 2.

To evaluate the associations of single-nucleotide polymorphisms (SNPs), LEPR Gln233Arg, LEP G2548A and BDNF Val66Met, with leptin level, BDNF level and metabolic features we performed multivariate analysis.

Table 1 Comparison of anthropometric, biochemical variables and age between individuals with and without MS

| Variables         | Control (n = 162) Median (range) | MS (n = 160) Median (range) | p-value |
|-------------------|----------------------------------|-----------------------------|---------|
| Age (years)       | 46.0 (24.0-64.0)                 | 48.0 (28.0-63.0)            | 0.241   |
| Leptin (ng/mL)    | 6.2 (2.2-32.3)                   | 11.4 (2.9-36.8)             | <0.001**|
| BDNF (ng/mL)      | 11.7 (2.8-29.2)                  | 11.5 (2.7-27.8)             | 0.946   |
| Glucose (mg/dL)   | 84.0 (66.0-113.0)                | 95.5 (70.0-134.0)           | <0.001**|
| Insulin (µU/mL)   | 11.7 (4.5-30.7)                  | 15.8 (5.6-37.3)             | <0.001**|
| TG (mg/dL)        | 84.0 (58.0-270.0)                | 169.0 (63.0-302.0)          | <0.001**|
| HDL-C (mg/dL)     | 50.0 (28.0-770)                  | 40.0 (27.0-710)             | <0.001**|
| Systolic BP (mmHg)| 120.0 (95.0-160.0)              | 130.0 (100.0-170.0)         | <0.001**|
| Diastolic BP (mmHg)| 74.0 (60.0-95.0)              | 80.0 (64.0-100.0)           | <0.001**|
| BMI (kg/m²)       | 22.8 (18.9-35.5)                 | 26.5 (19.5-37.9)            | <0.001**|
| Waist/Hip ratio   | 0.82 (0.69-0.97)                 | 0.88 (0.72-0.99)            | <0.001**|
| WC (cm)           | 77.0 (61.5-101.3)                | 91.0 (66.5-106.0)           | <0.001**|
| AC (cm)           | 27.5 (21.8-34.0)                 | 30.2 (22.9-38.5)            | <0.001**|
| TSF (mm)          | 17.9 (6.5-37.5)                  | 26.0 (8.5-40.9)             | <0.001**|

Table 2 Correlation coefficients of leptin as well as BDNF with other parameters, in all subjects

| Variables     | Leptin       | BDNF        |
|---------------|--------------|-------------|
| Leptin        | 1            | 0.021       |
| BDNF          | 0.021        | 1           |
| Age           | 0.233**      | -0.280**    |
| Glucose       | 0.097        | -0.059      |
| Insulin       | 0.152*       | -0.127      |
| TG            | 0.230**      | 0.127       |
| HDL-C         | -0.214**     | -0.057      |
| Systolic BP   | -0.358**     | -0.018      |
| Diastolic BP  | -0.199**     | -0.015      |
| BMI           | 0.383**      | 0.036       |
| Waist/Hip ratio | 0.301**   | 0.027       |
| WC            | 0.293**      | 0.052       |
| AC            | 0.408**      | 0.314**     |
| TSF           | 0.516**      | 0.224**     |

Significant associations of LEPR Gln223Arg polymorphism were found with leptin and glucose levels, after adjusting for potential covariates including age, sex, weight and smoking status; the results are shown in Table 3. There were no significant associations of LEP G2548A and BDNF Val66Met polymorphisms with leptin level, BDNF level and metabolic features (p > 0.05); the results are shown in Table 3. The genotypes of SNPs LEPR Gln233Arg, LEP G2548A and BDNF Val66Met were in Hardy-Weinberg equilibrium (HWE) (p > 0.05) and the values of chi-square and degrees of freedom were shown in Table 4. Moreover, the results in genotypic and allelic frequencies of LEPR Gln233Arg, LEP G2548A, and BDNF Val66Met polymorphisms in MS and control subjects are also shown in Table 4. There were significant differences in genotypic and allelic frequencies of LEPR Gln233Arg polymorphism between MS and control groups (p < 0.05), but genotypic and allelic distributions of LEP G2548A and BDNF Val66Met polymorphisms between these two groups were not significantly different (p > 0.05).

Association between LEPR Gln233Arg, LEP G2548A and BDNF Val66Met polymorphisms and MS among study subjects are shown in Table 5. Polymorphism of LEPR Gln233Arg showed a significant link to MS (OR = 1.8, p < 0.05), and Gln/Arg with Arg/Arg genotype frequencies in the MS and control groups were 62.5% and 48.8%, respectively. In contrast, LEP G2548A and BDNF Val66Met polymorphisms were not related to MS (p > 0.05).
### Table 3 Adjusted odds ratio for LEPR Gln223Arg, LEP G2548A and BDNF Val66Met polymorphisms with leptin level, BDNF level and metabolic features

| Variables          | SNP LEPR Gln223Arg | p-value | SNP LEP G2548A | p-value | SNP BDNF Val66Met | p-value |
|--------------------|---------------------|---------|----------------|---------|------------------|---------|
|                    | OR (95% CI)         |         | OR (95% CI)    |         | OR (95% CI)      |         |
| Systolic BP        | 0.989 (0.952-1.027) | 0.568   | 1.030 (0.982-1.090) | 0.318 | 1.022 (0.978-1.068) | 0.331 |
| Diastolic BP       | 1.019 (0.968-1.072) | 0.474   | 1.000 (0.932-1.073) | 0.999 | 1.017 (0.972-1.075) | 0.447 |
| WC                 | 0.991 (0.972-1.046) | 0.437   | 1.044 (0.966-1.129) | 0.270 | 0.968 (0.899-1.043) | 0.410 |
| TG                 | 0.999 (0.993-1.004) | 0.635   | 1.002 (0.994-1.009) | 0.636 | 0.998 (0.991-1.007) | 0.378 |
| HDL-C              | 1.000 (0.974-1.038) | 0.994   | 0.992 (0.945-1.041) | 0.745 | 1.002 (0.965-1.029) | 0.722 |
| Glucose            | 1.047 (1.004-1.092) | 0.033*  | 1.021 (0.989-1.065) | 0.340 | 0.989 (0.965-1.019) | 0.354 |
| Insulin            | 0.916 (0.997-1.056) | 0.999   | 0.985 (0.944-1.027) | 0.473 | 1.009 (0.972-1.050) | 0.600 |
| Leptin             | 1.192 (1.038-1.379) | 0.009** | 0.924 (0.816-1.039) | 0.209 | 0.963 (0.873-1.050) | 0.903 |

**Abbreviation:** BDNF Brain-derived neurotrophic factor, TG triglyceride, HDL-C high-density lipoprotein cholesterol. WC waist circumference, LEP leptin gene, LEPR leptin receptor gene, BDNF Brain-derived neurotrophic factor gene.

Significant levels: * = p < 0.05, **= p < 0.01.

### Table 4 Genotypic and allelic distribution of LEPR Gln233Arg, LEP G2548A, and BDNF Val66Met polymorphisms in MS and control subjects

| Genotype | MSn (% | HWE of MS p-value | Control n (%) | HWE of control p-value | Genotypic or allelic p-value |
|----------|--------|-------------------|----------------|------------------------|-----------------------------|
| Arg/Arg  | 20 (12.5) | 0.399 | 17 (10.5) | 0.295 | | |
| Arg/Gln  | 80 (50.0) | (df = 1) | 62 (38.3) | (df = 1) | | 0.044* |
| Gln/Gln  | 60 (37.5) | (χ² = 0.71) | 83 (51.2) | (χ² = 1.09) | | |
| Allele   | Arg 120 (0.375) | 96 (0.296) | | | 0.034* |
|          | Gln 200 (0.625) | 228 (0.704) | | | |

**LEP G2548A**

| Genotype | MSn (% | HWE of MS p-value | Control n (%) | HWE of control p-value | Genotypic or allelic p-value |
|----------|--------|-------------------|----------------|------------------------|-----------------------------|
| A/A      | 58 (36.2) | 0.638 | 55 (34.0) | 0.483 | | |
| A/G      | 79 (49.4) | (df = 1) | 75 (46.3) | (df = 1) | | 0.439 |
| G/G      | 23 (14.4) | (χ² = 0.22) | 32 (19.7) | (χ² = 0.49) | | |
| Allele   | A 195 (0.609) | 185 (0.571) | | | 0.321 |
|          | G 125 (0.391) | 139 (0.429) | | | |

**BDNF Val66Met**

| Genotype | MSn (% | HWE of MS p-value | Control n (%) | HWE of control p-value | Genotypic or allelic p-value |
|----------|--------|-------------------|----------------|------------------------|-----------------------------|
| Met/Met  | 32 (20.0) | 0.479 | 30 (18.5) | 0.544 | | |
| Met/Val  | 84 (52.5) | (df = 1) | 75 (46.3) | (df = 1) | | 0.327 |
| Val/Val  | 44 (27.5) | (χ² = 0.50) | 57 (35.2) | (χ² = 0.37) | | |
| Allele   | Met 148 (0.463) | 135 (0.417) | | | 0.241 |
|          | Val 172 (0.537) | 189 (0.583) | | | |

**Abbreviation:** HWE Hardy-Weinberg equilibrium, LEP leptin gene, LEPR leptin receptor gene, BDNF Brain-derived neurotrophic factor gene.

Significant levels: * = p < 0.05, **= p < 0.01.

a Based on the results of chi-square test.
b Based on the results of chi-square test for the comparison between MS and control groups.
Logistic regression results for possible associations between MS and age, sex, TSF, waist/hip ratio, insulin, leptin, LEP G2548A, LEPR Gln223Arg, and BDNF Val66Met, are shown in Table 6. Of these variables, leptin, LEPR Gln223Arg, TSF and waist/hip ratio were significantly related to MS (p < 0.05). The Hosmer-Lemeshow Goodness-of-Fit test ($\chi^2 = 0.896$, p > 0.05) was not statistically significant; the fit between the predictive model and the data was acceptable.

### Discussion

To the best of our knowledge, this is the first study to examine the links of leptin level, BDNF level, and polymorphisms of LEPR Gln223Arg, LEP G2548A and BDNF Val66Met with MS among Thais. The key observations of this study were that leptin levels were significantly higher in the MS group than in the control group, and that the LEPR Gln223Arg polymorphism showed a significant link to leptin concentrations and MS among Thais. In contrast, LEP G2548A and BDNF Val66Met polymorphisms were not related to MS.

MS is a cluster of risk factors predictive of future cardiovascular diseases and type 2 diabetes mellitus [1]. Factors which appear to promote the development of MS include genetics, obesity, physical inactivity and an unhealthy diet. Mechanisms causing the onset of this syndrome, however, are still not clearly understood. Leptin is an adipocyte-derived signaling factor that has an important role in metabolic control, such as in stimulating glucose uptake, fatty acid oxidation, and reducing food intake [2,22]. Moreover, circulating leptin levels are closely related to stored adipose tissues [3] and metabolic function [22]. However, most previous studies have focused on the relationship between leptin and obesity, and each component of MS, instead of a cluster of MS components. Therefore, the present study investigated links between leptin and MS among Thais, especially. Our results were in accordance with previous studies in Korean populations [23] and Lebanese volunteers [24], in that serum leptin was associated with MS. Bremer et al. also showed increased amounts of subcutaneous adipose tissue secretion and circulating leptin concentrations in subjects with nascent MS [25]. Increased leptin concentration in MS may represent a form of leptin resistance, as in obese subjects [26]; obesity is one of the risk factors for MS. Therefore, alternation of leptin levels may be a part of the etiology of MS.

Leptin exerts a number of significant influences, by binding and activating specific leptin receptors in the hypothalamus and other organs including pancreatic β cells, adipose tissue, and muscle. Previous studies reported LEPR polymorphisms were associated with glucose metabolism [27] and insulin resistance [5], possibly leading to a predisposition towards greater MS risk. LEPR Gln223Arg polymorphism has been associated with impaired signaling capacity of the leptin receptor [8]. LEPR Gln223Arg polymorphism has also been linked with impaired glucose tolerance and conversion to type 2 diabetes [27]. Our study was therefore interested in this position of the LEPR gene polymorphism. Present data points to a relationship between LEPR Gln223Arg polymorphism and increased leptin and glucose levels in Thai subjects, similar to results in Dutch adults [28] and Romanian subjects [29]. In contrast to our results, however, LEPR Gln223Arg polymorphism

### Table 5 Association between LEPR Gln233Arg, LEP G2548A and BDNF Val66Met polymorphisms and MS

| SNPs        | Genotype frequencies | OR (95% CI) | p-value* |
|-------------|----------------------|-------------|----------|
| LEPR Gln223Arg |                      |             |          |
| Arg/Arg + Arg/Gln | 100 (62.5) 79 (48.8) | 1.8 (1.2-2.7) | 0.013*   |
| Gln/Gln     | 60 (37.5) 83 (51.2)  |             |          |
| LEP G2548A  |                      |             |          |
| A/A + A/G   | 137 (85.6) 130 (80.2) | 1.5 (0.8-2.6) | 0.200    |
| G/G         | 23 (14.4) 32 (19.8)  |             |          |
| BDNF Val66Met |                    |             |          |
| Met/Met + Met/Val | 116 (72.5) 105 (64.8) | 1.4 (0.8-2.5) | 0.230    |
| Val/Val     | 44 (27.5) 57 (35.2)  |             |          |

Abbreviation: LEP: leptin gene, LEPR: leptin receptor gene, BDNF: brain-derived neurotrophic factor gene. Significant level* = p < 0.05.

### Table 6 Logistic regression analysis when MS was used as a dependent variable, and age, sex, TSF, waist/hip ratio, insulin, leptin, LEPR Gln223Arg, LEP G2548A and BDNF Val66Met, were taken as independent variables

| Variables | β     | p-value | OR (95% CI) |
|-----------|-------|---------|-------------|
| Leptin    | 0.46  | 0.006*  | 1.91 1.25-3.03 |
| LEPR Gln223Arg | 2.37  | 0.036* | 9.07 1.22-3.14 |
| Waist/hip ratio | 5.58  | <0.001* | 22.04 2.42-91.36 |
| TSF       | 0.17  | 0.022*  | 1.31 1.03-1.95 |
| Sex       | -2.34 | 0.331   | 0.20 0.02-6.26 |
| Age       | 0.00  | 0.171   | 1.01 0.09-11.0 |
| Insulin   | 0.07  | 0.010*  | 1.11 0.98-1.21 |
| BDNF Val66Met | 0.88  | 0.117   | 2.39 0.74-8.54 |
| LEP G2548A | 1.84  | 0.124   | 6.03 0.80-26.11 |

Abbreviation: LEP: leptin gene, LEPR: leptin receptor gene, BDNF: brain-derived neurotrophic factor gene. Significant level* = p < 0.05.
demonstrated no relationship with leptin levels in Turkish children [11]. Moreover, few studies have determined a correlation between LEPR gene polymorphism and MS. Our findings among Thais were in accordance with results from a sample of elderly Brazilian subjects [9], which showed links between LEPR Gln223Arg polymorphism and MS. On the other hand, the study in Turkish children did not find any relation between LEPR Gln223Arg polymorphism and MS [11]. Therefore, our study demonstrated that LEPR Gln223Arg polymorphism in Thai subjects was associated with increased MS risk, which may be explained by increasing glucose and leptin levels. Although the pathogenesis of MS and each of its components is complex and not well understood, our investigation proposes that LEPR Gln223Arg polymorphism associated with increased leptin levels may represent a modified functional leptin receptor and may take part in the development of MS. The previous studies also suggested that LEPR Gln223Arg polymorphism has been associated with insulin resistance capacity and an altered leptin-binding activity [30] and the leptin can modulates insulin secretion and action via leptin receptors in pancreatic β cells [31].

Although many researches indicate a role for genetic susceptibility to the MS and our results supporting the relationship between the LEPR Gln223Arg polymorphism and increased MS risk among Thais, lifestyle factors have also found to be affecting factors in the pathogenesis and progression of MS. Thailand has importantly experienced economic transition and its structure has gradually changed from the traditional agricultural surroundings to an industrialized structure. The dietary intake pattern in Thais has changed from traditional high-carbohydrate diets, which rely heavily on rice and vegetables, to diets high in fat and sugar; physical activity has progressively declined [32]. These transitions have consequently resulted in increasing the prevalence of MS in Thailand [33]. Therefore, lifestyle interventions including eating healthy diet and improving physical activity should be regarded as important management for reducing the development of MS among Thais, especially in SNP LEPR Gln223Arg subjects. With regard to links between LEP G2548A gene polymorphism and leptin levels, the previous study reported a common gene polymorphism in the promoter LEP G2548A influencing leptin expression and adipose tissue secretion of leptin [7]. However, studies yielded inconsistent results; data in Egyptian subjects [34] and Romanian subjects [29] points to a relationship between LEP 2548GG variants and serum leptin levels, meanwhile studies in Melanesian and Micronesian Solomon Islanders [35] found no relationship between LEP G2548A polymorphism and leptin levels, consistent with our results among Thais. In investigating the links between LEP G2548A polymorphism and MS, LEP G2548A polymorphism was shown to be significantly associated with MS in Tunisian volunteers [10] and Egyptian subjects [34], while no links were found in our study. Investigations among Romanian subjects also found no association of the LEP G2548A polymorphism with common obesity-related variables and metabolic traits [29]. It is possible that inconsistencies among the results of the LEP and LEPR gene polymorphisms in these studies are the result of the different genetic backgrounds or environmental conditions of the populations studied.

BDNF is involved in regulating energy homoeostasis, blood glucose levels, and eating behaviors, and it appears to function downstream of the leptin-melanocortin signaling pathway [13,36]. BDNF is expressed at high levels in the ventromedial hypothalamus (VMH) where its expression is regulated by nutritional state and by melanocortin-4 receptor (MC4R) signaling [36]. Moreover, Komori et al. demonstrated that administration of leptin causes an increase in expression of BDNF mRNA and protein in the VMH [37]. The precise mechanism that mediates BDNF expression by leptin remains to be elucidated and there are two evidences that have been proposed. The first possibility is that the LEP-LEPR interaction directly triggers a signal transduction cascade that induces BDNF in the VMH. The second option involves the leptin-mediated production of the melanocortin precursor α-melanocyte stimulating hormone (a-MSH) in the arcuate nucleus, which activates BDNF in the VMH via MC4R [13]. However, the mechanistic implication of the association between leptin, BDNF and the LEPR gene in MS is unclear. Our results may imply that Thai MS subjects have related to LEPR Gln223Arg polymorphism and increased leptin levels representing a form of leptin resistant. These defects may be involved in the LEP-LEPR interaction in mechanisms induced BDNF expression by leptin signaling. Although previous studies have investigated a correlation between BDNF levels and MS, these investigations have shown inconsistent results of BDNF levels in subjects with MS [16,17,38,39]. In examining links between BDNF concentrations and BDNF Val66Met gene polymorphisms with MS among Thais, the present study found neither BDNF concentrations nor BDNF gene polymorphism relates to MS. Our results were in line with studies in Spanish and Taiwanese subjects, in that, BDNF concentrations did not correlate with MS and any metabolic syndrome component [16,38]. In contrast, other reports have found BDNF levels to be linked with MS, and with several features of MS [17,39]. However, data on BDNF Val66Met gene polymorphism with MS is scarcely available. Indeed, ours is the first publication to determine these associations among Thais. Previous studies found that BDNF Val66Met gene polymorphism influenced BDNF expression and/or activity [15], and was related to...
a higher risk of becoming obese [40]. One study among Caucasian subjects found no significant association between Val66Met gene polymorphism and MS [41]. Similarly, our results confirm this polymorphism not to be linked with MS or with features of MS. Therefore, our findings suggest that BDNF concentrations and BDNF Val66Met polymorphism are not involved in the pathophysiology of MS among Thais. However, the relationship between BDNF levels and gene polymorphisms with MS is still not clearly understood, and further studies should be undertaken investigating different ethnicities, as well as other points of the BDNF gene.

There are a few limitations of our study. We did not analyze the whole gene of LEP, LEPR and BDNF with MS and further studies should be replicated our findings in larger sample numbers.

In conclusion, the present study found leptin concentrations of MS subjects to be higher than the controls. Although no correlations between MS with BDNF level or BDNF Val66Met were observed, there was an association between LEPR Gln223Arg polymorphism and leptin level and MS. Therefore, genetic polymorphism in the LEPR Gln223Arg appeared to affect the leptin concentration and the susceptibility to MS and these findings might play a significant function in the etiology of MS among the Thai population. Moreover, we believe further studies on MS genetics will help to develop preventive strategies and the role of other genetic and environmental factors should be studied in the other populations.

Competing interests
The authors have no conflicts of interest to declare.

Authors' contributions
KS contributed to the conception and study design, collected specimens, carried out laboratory analysis and prepared the manuscript. KT carried out laboratory analysis and helped to draft the manuscript. RT collected specimens and interpretation of the data. All authors read and approved the final manuscript.

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