Abstract. The polycystic ovary syndrome (PCOS) is considered as one of the most common endocrine and reproductive dysfunctional diseases. Recent research for genetic variants has identified genetic influences between the polymorphisms of the adiponectin gene and the metabolic syndromes. The aim of our study was to investigate the relationship between single nucleotide polymorphisms (SNPs) in the adiponectin gene and PCOS. Two SNPs, +45G15G(T/G) and +276(G/T), which are found in exon 2 and intron 2, respectively, of the adiponectin gene, were genotyped by PCR-RFLP. Out of 303 women studied for the +45G15G(T/G) and +276(G/T) SNPs, 144 had PCOS and 159 were healthy controls. No association was found between the +45(T/G) SNP and PCOS (P=0.3558, OR=0.83, 95% confidence interval), per contra to the association between +276(G/T) SNP and PCOS (P=0.0126, OR=0.60, 95% confidence interval). These results indicate that the SNP of +276(G/T) is strongly associated with PCOS. However, the +45(T/G) SNP is not associated with PCOS.

Introduction

The polycystic ovary syndrome (PCOS) is a common multifactorial disease, also considered as an endocrine syndrome with polymorphic clinical manifestations, including hyperandrogenism and chronic anovulation (1-5). The current diagnostic criteria are based on the 2003 American Society for Reproductive Medicine/European Society of Human Reproduction and Embryology (ASRM/ESHRE) Rotterdam consensus (6,7). So far, the aetiology of PCOS has not been fully elucidated, but correlation studies have found that PCOS is likely to be hereditary, since a sister with PCOS increases the risk of PCOS up to 46% (8). Therefore, the study of the PCOS candidate gene will reveal the fundamental mechanism and complication of PCOS.

A large number of studies have shown that there is a close relationship between insulin resistance (IR) and PCOS. IR is a main factor in the development of PCOS, and about 64.4% of PCOS patients have IR (9). PCOS patients, whether obese or not, have different levels of IR and hyperinsulinemia and IR is independent of obesity. IR leads to obstacles in fat mobilization and utilization, becoming an important risk factor for obesity. Furthermore, lipocytes, one of the major insulin target cells, can produce a variety of factors involved in the development of IR (10,11).

Adiponectin is one of the most abundant adipokines and accounts for 0.01% or 3-30 μg/ml of the total plasma protein (12). Adiponectin is secreted by lipocytes and is composed of 224 amino acid peptides. Several studies have shown that adiponectin plays an important role in the process of glucose regulation and lipid metabolism. Moreover, adiponectin can increase skeletal muscle fatty acid oxidation and peripheral tissue sensitivity to insulin and can inhibit hepatic gluconeogenesis. It has been suggested that adiponectin has a protective effect on the development of atherosclerosis (13,14). In different animal models for obesity and diabetes, adiponectin was found to correlate with the insulin sensitivity index (15). Furthermore, adiponectin has been shown to be an important inflammatory response regulator (16).

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These include the following single nucleotide polymorphisms (SNPs): -11426(A/G), -11391(A/G), -11377(C/G), +45(T/G), +276(G/T) and H111Y(C/T) (11-12,17-21). Intriguingly, SNPs +45(T/G) and +276(G/T) which are located in exon 2 and intron 2 of the adiponectin gene, respectively, have been found to be strongly associated with obesity, IR and T2D in Asian populations. In this study, we assessed the association of SNPs +45(T/G) and +276(G/T) in the adiponectin gene with PCOS in the Korean population.

Materials and methods

Subjects. All of the PCOS patients and controls were Korean women which were recruited from the Fertility Center of the CHA General Hospital in Seoul, Korea. Informed consent was obtained from all patients. Among 303 Korean women, 144 had PCOS, while the remaining 159 were healthy control subjects. The diagnosis of PCOS was based on the criteria proposed by the 2003 ASRM/ESHRE Rotterdam consensus. Blood samples were collected in tubes containing EDTA as an anticoagulant and stored at 4°C. Genomic DNA was extracted from the blood of PCOS patients and control women. The study was approved by the Institutional Review Board.

Biochemical determinations. Blood samples were obtained from PCOS patients and control subjects for chemical analyses of FSH, LH, TSH, prolactin, DHEA-S, and E2 as indicators of discrimination.

Genetic analysis. Two SNPs were genotyped by the PCR-based RFLP analysis for all subjects. Primers and restriction enzymes used are shown in Table I. The PCR amplification conditions used for +45G15G(T/G) were as follows: 94°C for 5 min, followed by 30 cycles of 35 sec at 94°C, 30 sec at 58.8°C, and 30 sec at 72°C. PCR conditions for +276G(T/T) were as follows: 95°C for 5 min, followed by 35 cycles of 30 sec at 95°C, 30 sec at 62°C, and 30 sec at 72°C. The PCR products were purified using Bioneer’s AccuPrep PCR purification kit (Bioneer, Daejeon, Korea) and were digested with SmaI (New England Biolabs, Beverly, MA, USA) for 6 h at 25°C or BsmI (New England Biolabs) for 14 h at 65°C for SNPs +45(T/G) and +276(G/T), respectively. The digested DNA fragments were electrophoresed on a 2% agarose gels containing ethidium bromide and visualized by an ultraviolet transilluminator. In regard to the SNP +45(T/G), a single 390-bp band indicates homozygosity for the T allele; the two bands, 217- and 173-bp, indicate homozygosity for the G allele; the presence of three fragments, 390-, 217- and 173-bp bands, indicate heterozygosity for the T or G allele (Fig. 1). As for the SNP +276(G/T), presence of a single 468-bp band indicates homozygosity for the T allele; presence of two fragments, 320- and 148-bp, indicates homozygosity for the G allele; and three fragments, 468-, 320- and 148-bp indicate heterozygosity for the G and the T allele (Fig. 2).

Statistical analysis. Genotype and allele frequencies were compared between patients and controls by the \( \chi^2 \) test. Analyzed haplotypes composed of SNPs were evaluated with HapAnalyzer. P-values of <0.05 were considered as statistically significant.
Results

The analysis of the clinical and biochemical characteristics of the healthy control subjects and of patients with PCOS are shown in Table II. In regard to the diagnostic criteria of PCOS, we followed the 2003 ASRM/ESHRE Rotterdam consensus (6). According to the criteria, patients are diagnosed with PCOS when they have two of the following three features: oligo- or amenorrhea, clinical or biochemical hyperandrogenism and ultrasonographic polycystic ovarian morphology (6). As shown in Table II, there was difference in the level of TSH, DHEA-S and testosterone for the PCOS patients and control groups. In addition, the level of LH in PCOS patients was almost twice as much as in the control group. In the PCOS patient group, 22 patients (15.30%) had hyperandrogenism and oligo- or amenorrhea, 12 patients (8.33%) had hyperandrogenism and polycystic ovaries, 94 patients (65.28%) had oligo- or amenorrhea and polycystic ovaries, and 15 patients (10.42%) had hyperandrogenism, oligo- or amenorrhea and polycystic ovaries (Table II).

To analyze the frequency of the genotypes for SNPs in the adiponectin gene, we performed PCR-RFLP analysis. Table III shows the SNP allele frequency of the TT, TG, GG genotypes of the exon 2 T/G polymorphism and of the GG, GT, TT genotypes of the intron 2 G/T polymorphism for the adiponectin gene in patients with PCOS and control subjects. In the case of +45 (T/G), the frequency of the TT genotype was slightly higher in the PCOS patient group than in the control group (54.9% vs. 45.3%, respectively); the TG genotype was more frequent in the control group (52.8%) than in the PCOS patient group (41.0%); the frequency of the GG genotype was similar between the two groups (patient group, 4.2%; control group, 1.9%). Also, for the +276 (G/T) SNP, significantly higher frequency of the GG genotype was shown in the PCOS patient group than in the control group (42.4% vs. 30.2%, respectively); the rate of the GT genotype was slightly higher in the control group than in the PCOS patient group; and the frequency of the TT genotype was remarkably higher in the control group (15.1%) than in the PCOS patient group (6.9%). In addition, as indicated in Table III, there was no association between the SNP +45 (T/G) and the occurrence of PCOS. In contrast, the data demonstrate the association between the SNP +276 (G/T) and PCOS patients in Korean women.

Discussion

Molecular genetic studies have shown that hormone metabolism and energy regulation-related genes, such as CYP19, INSR, LH are involved in the pathogenesis of PCOS (22-26).
In the present study, we assessed another metabolism-related gene, adiponectin, and examined the association between the SNPs +45(G/T) and +276(G/T) in the adiponectin gene and PCOS in Korean women. An association was found between the +276(G/T) SNP and PCOS, while the +45(T/G) SNP was not associated with PCOS in Korean women. To a certain extent, variation of the adiponectin gene determines the overexpression of the phenotype of the metabolic syndrome. However, it has been reported that the polymorphism of the adiponectin gene is not the real reason for the metabolic disorders of PCOS (27); it has been suggested that a certain degree of interaction may exist between adiponectin and steroid hormone activity and synthesis. At present, 13 SNPs have been identified in the adiponectin gene (28), and most of the studies have focused on two SNPs: +45(T/G) and +276(G/T). However, the influence of these two polymorphisms on the adiponectin gene expression or biological function is not fully known, because the SNP +45(T/G) is a synonymous polymorphism while the SNP +276(G/T) is an intronic one.

It has been reported that SNPs +45(T/G) and +276(G/T) in the adiponectin gene are significantly associated with PCOS in Chinese women (29). These two SNPs have been analyzed in various diseases and in different ethnic backgrounds. It has been reported that fasting glucose levels are influenced by the adiponectin +276(G/T) polymorphism in Korean and Italian populations (30,31). In French and Swedish Caucasians, the +45(T/G) and +276(G/T) polymorphisms have not been associated with T2D (32). However, it has also been indicated that SNPs +45(T/G) and +276(G/T) are strongly associated with type 1 diabetes (T1D), but not with diabetic nephropathy (DN) in Swedish patients (33). Studies have illustrated that the SNP in intron 2 of the adiponectin gene plays an important role in the regulation of adiponectin levels, although the exact cellular mechanism is still not known (34,35).

This is the first report on the association of two SNPs of the adiponectin gene with PCOS patients in Korea. Our results provide insight into the role of the adiponectin gene in the pathogenesis of PCOS in different ethnic backgrounds and also provide an important basis for the early diagnosis and prevention of PCOS.

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