

# Association study between the -866G/A polymorphism in the promoter of *uncoupling protein-2* gene and polycystic ovary syndrome

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**Abstract.** Polycystic ovary syndrome (PCOS) is a disorder characterized by hyperandrogenism, chronic oligoanovulation and insulin resistance. A number of women with PCOS are obese and exhibit abnormal phenotypes, including high levels of androgens, an irregular menstrual cycle and increased hair growth. Studies on obese PCOS patients have proven the crucial role that obesity plays in insulin resistance and hyperinsulinemia. The *uncoupling protein (UCP)* gene is one of the genes known to have a strong association with obesity and insulin resistance. Thus, we analyzed the association between the -866G/A polymorphism in the promoter of *UCP2* and PCOS. Genotyping was performed by polymerase chain reaction along with restriction fragment length polymorphism analysis, followed by an analysis of the genotype of the *UCP2* polymorphism in PCOS and control subjects using HapAnalyzer. The study included samples from 277 PCOS patients and 152 healthy controls.  $P < 0.05$  was considered to be statistically significant. In conclusion, no association was found between the -866G/A single nucleotide polymorphism and PCOS ( $P = 0.7168$ ,  $OR = 1.07$ , 95% CI). The present study showed that -866G/A, a *UCP2* gene polymorphism, is not associated with the pathogenesis of PCOS.

## Introduction

Polycystic ovary syndrome (PCOS), one of the most common endocrine disorders, affects 5-10% of women. It is characterized by hyperandrogenism and chronic oligoanovulation (1-6). PCOS patients are reported to have high serum concentrations of androgenic hormones, including testosterone, androstenedione and dehydroepiandrosterone sulfate (DHEA-S) (1-5).

PCOS is also associated with the metabolic syndrome, which is linked to insulin resistance, hyperinsulinemia and obesity. The post-binding defect in the insulin receptor signaling pathway may be one of the reasons for insulin resistance in PCOS, while the high insulin level potentially has a gonadotropin-augmenting effect on ovarian function (3,5,7). Obesity is also a significant common clinical feature in women affected by PCOS. Over 50% of patients were found to be overweight or obese (5,8), which may occur due to the onset of oligomenorrhea and hyperandrogenism, suggesting a pathogenic role of obesity in the development of the syndrome (5).

Uncoupling proteins (UCPs) are mitochondrial carrier proteins that uncouple the transport of protons across the inner mitochondrial membrane from electron transport and are involved in the synthesis of ATP from ADP (9-10). The archetypal *UCP1* is mainly expressed in brown adipocytes, while *UCP2* expression is widely distributed in mammalian tissues, including white adipose tissue, skeletal muscle, pancreatic islets, and the central nervous system (11-13). A number of polymorphisms and mutations in three uncoupling protein homolog genes (*UCP1*, *UCP2* and *UCP3*) have been reported (10). The polymorphisms in these *UCPs* have been shown to play a significant role in the pathogenesis of type 2 diabetes (T2D) mellitus and obesity (10). Perturbation in the expression level of *UCP2* or altered action of *UCP2* leads to T2D, disordered lipid metabolism, impaired insulin secretion and dysfunctional weight homeostasis (10,14-16).

There are three main reported polymorphisms in the *UCP2* gene that are associated with changes in body mass index (BMI), energy expenditure and maintenance of body weight after overfeeding (10,17). Various studies have been performed on exon 4 of the *UCP2* gene for the substitution of valine (V) for alanine (A) (Ala55Val) or the 45-bp insertion/deletion variant (10,18). If not all, the majority of these polymorphism studies on the *UCP2* gene revealed an association with obesity and energy expenditure. Esterbauer *et al* identified a functional polymorphism located in the promoter region of the *UCP2* gene (17). This polymorphism contributes to 71% of variation in the expression ratio between the insertion and deletion alleles (17). The -866A allele has been shown to enhance or suppress *UCP2* transcriptional activity in transfected cultured

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cells (13,19), indicating that *UCP2* mRNA levels are associated with the -866A allele. Additionally, the -866G allele is associated with an increased risk of chronic inflammatory diseases (20), and susceptibility to autoimmune (21) and cardiovascular diseases (22).

A number of PCOS patients with obesity exhibit the conditions of insulin resistance and hyperinsulinemia. Previous studies demonstrated the mechanism by which obesity may induce an insulin-resistant state in PCOS patients. Additionally, these studies emphasized the role of obesity in amplifying the degree of hyperandrogenism in PCOS (5,23). A growing body of evidence reports that the -866G/A polymorphism of *UCP2* is associated with T2D and obesity (10,12), which led to our hypothesis that the -866G/A polymorphism of *UCP2* plays a significant role in obese PCOS patients. The purpose of the present study was to analyze the -866G/A polymorphism in the *UCP2* promoter in obese PCOS patients. This was the first study to investigate the association between the -866G/A polymorphism in the promoter of *UCP2* and PCOS.

## Materials and methods

**Subjects.** A total of 429 Korean females, of which 152 were healthy control subjects and 277 were PCOS patients, were selected from the Fertility Center at CHA General Hospital in Seoul, Korea, to participate in this study. The diagnosis of PCOS was based on the criteria proposed by the 2003 ASRM/ESHRE Rotterdam consensus. This study was approved by the Institutional Review Board (IRB).

**DNA extraction and genotyping.** For molecular genetic experiments, blood samples from the PCOS patients and the control group were collected in tubes containing EDTA as an anticoagulant and were stored at 4°C. Genomic DNA was then extracted from the whole blood samples.

The -866G/A variant of *UCP2* was amplified using a forward primer: 5'-CAC GCT GCT TCT GCC AGG AC-3' and a reverse primer: 5'-AGG CGT CAG GAG ATG GAC CG-3'. Cycling parameters were as follows: denaturation at 94°C for 5 min, 30 cycles at 94°C for 30 sec, annealing at 65°C for 30 sec, extension at 72°C for 30 sec and a final at 72°C for 5 min. The PCR products of 360-bp were purified using Accuprep Bioneer's PCR purification kit (Bioneer, Daejeon, Korea), and digested with *MluI* (New England Biolabs, Beverly, MA, USA) (Fig. 1A). The digested fragments were analyzed on 2% agarose gel. Upon digestion with *MluI*, the A allele was cleaved into two fragments (290- and 70-bp), whereas the G allele was identified by a single band (360-bp) (Fig. 1B).

**Biochemical determinations.** Blood samples from the PCOS patients and controls were analyzed using biochemical assays, including FSH, LH, TSH, PRL, DHEA-S and E2.

**Statistical analysis.** Genotype frequencies of the patients and control samples were compared using the  $\chi^2$  test, and analyzed using HapAnalyzer (NGRI, Seoul, Korea; www.hap.ngri.re.kr).  $P < 0.05$  was considered to be statistically significant. To verify whether the haplotypes in the -866G/A polymorphism of *UCP2* were associated with PCOS, the results were compared using HapAnalyzer.

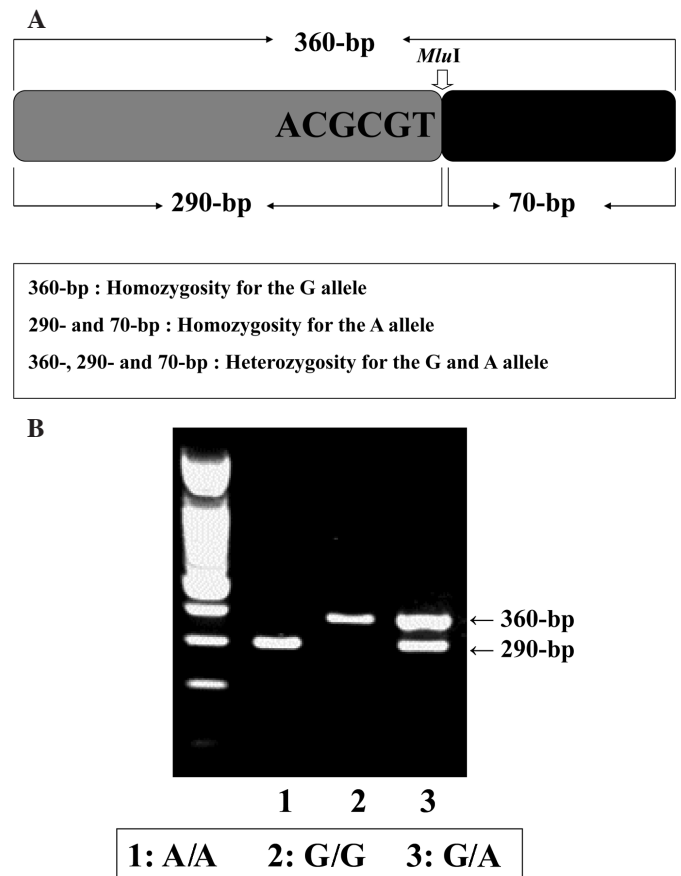


Figure 1. (A) Structure of the *UCP2* gene. The arrow indicates the restriction site of the *UCP2* gene by *MluI*. *MluI* restricts the site and produces two fragments, 290- and 70-bp in the sequence containing the A allele. The sequence containing the G allele yields only one fragment of 360-bp. (B) G/A polymorphism of the *UCP2* gene. The A allele is restricted by *MluI*. Agarose gel electrophoresis (2%) with ethidium bromide staining followed *MluI* digestion of the PCR product. The single band shows 360-bp of the G/G genotype. Homozygosity of A allele has two bands of 290- and 70-bp. However, the 70-bp band is not detected on the 2% agarose gel. Three bands of 360-, 290- and 70-bp indicate the G/A type.

## Results

The 2003 ASRM/ESHRE Rotterdam consensus was adhered to for the diagnostic criteria of PCOS in order to recruit patients into the study (24-25). According to this standard, PCOS may be diagnosed when patients exhibit two of three phenotypes, i.e., oligo- or amenorrhea, clinical or biochemical hyperandrogenism and ultrasonographic polycystic ovarian morphology (1,3,4). The analysis of clinical and biochemical characteristics revealed a significant difference between the PCOS and control groups in the levels of LH, TSH, DHEA-S and testosterone. The LH level in PCOS patients was found to be 2-fold higher than that in the control group (Table I). The results obtained from the biochemical analysis on the PCOS patient group revealed that 55 patients (19.86%) were susceptible to hyperandrogenism and oligo- or amenorrhea, 45 (16.24%) had hyperandrogenism and polycystic ovaries (PCOS), 129 (46.57%) had oligo- or amenorrhea and polycystic ovaries, and 48 (17.33%) had hyperandrogenism, oligo- or amenorrhea and polycystic ovaries. These data are recorded in Table II.

Table I. Biochemical and clinical characteristics of normal controls and PCOS patients.

Characteristics	Controls (n=152)	PCOS patients (n=277)
BMI (kg/m <sup>2</sup> )	20.75±2.39 (16.38-38.32)	22.49±2.48 (16.88-29.68)
Waist/hip ratio (WHR)	0.85±0.078 (0.73-0.95)	0.89±0.02 (0.71-1.15)
FSH levels (mIU/ml)	6.89±1.88 (3.29-9.79)	5.17±1.19 (2.86-18.58)
LH levels (mIU/ml)	3.28±1.66 (2.51-11.55)	7.00±5.88 (1.00-21.00)
E2 levels (pg/ml)	34.97±27.88 (4.79-218.09)	31.79±18.45 (8.00-86.50)
Prolactin levels (ng/ml)	14.42±6.48 (5.10-42.60)	12.64±9.98 (6.30-68.30)
TSH levels (μIU/ml)	1.88±0.98 (0.05-4.08)	2.42±1.28 (0.56-12.56)
DHEA-S levels (μg/dl)	142.46±35.26 (56.63-236.20)	187.04±54.54 (38.00-350.20)
Testosterone (ng/ml)	0.23±0.14 (0.03-0.54)	0.47±0.24 (0.15-0.89)

BMI, body mass index.

Table II. Comparison of disorders/symptoms between the control group and PCOS patients.

Characteristics	Controls (n=152) (0%)	PCOS patients (n=277) (%)
Hyperandrogenism and oligo- or amenorrhea	n=0	n=55 (19.86)
Hyperandrogenism and polycystic ovaries	n=0	n=45 (16.24)
Oligo- or amenorrhea and polycystic ovaries	n=0	n=129 (46.57)
Hyperandrogenism, oligo- or amenorrhea and polycystic ovaries	n=0	n=48 (17.33)

Table III. Genotypes and allele frequencies of the -866G/A polymorphism of the *UCP2* gene in women with PCOS and the controls group.

Polymorphism	PCOS (n=277)	Controls (n=152)	P-value
-866G/A			
Genotypes, no. (%)			
A/A	57 (20.6)	38 (25.0)	0.7168
G/A	157 (56.7)	68 (44.7)	
G/G	63 (22.7)	46 (30.3)	
Alleles, no. (%)			
A	271 (48.9)	144 (47.4)	
G	283 (51.1)	160 (52.6)	

In order to perform genotypic analysis of the -866G/A polymorphism in the promoter of the *UCP2* gene, PCR-RFLP analysis was carried out on the 277 PCOS patients and

152 controls. As a result, the frequency of A/A, G/A and G/G genotypes was revealed to be in similar proportion in both the PCOS and control groups (Table III). The frequency of the A/A genotype in the PCOS and control groups was 20.6 vs. 25%, respectively; the frequency of the G/A genotype in the PCOS and control groups was 56.7 vs. 44.7%, respectively; and the frequency of the G/G genotype in the PCOS and control groups was 22.7 vs. 30.3%, respectively. These results reveal that there is no significant difference between the PCOS and control groups, indicating that no association exists between the -866G/A polymorphism in the promoter of the *UCP2* gene and PCOS.

### Discussion

UCP plays a key role in the regulation of human energy metabolism by dissipating proton gradients, uncoupling respiration from oxidative phosphorylation, and converting fuel to heat (26). In numerous studies, -866G/A, -55C/T and 45-bp insertion/deletion of the gene polymorphism were reported for *UCP2* (10,18). Among them, -866G/A SNP was identified in the promoter region (17). Notably, this variant is reportedly associated with fat metabolism, obesity and diabetes in the majority of the population studied, including a Korean population (18). The minor A allele variant was frequently reported to be associated with increased adipose mRNA expression *in vivo* and with a modest reduction in obesity (17,27). The -866A allele carrier is generally susceptible to developing diabetes due to a reduced insulin response to intravenous and oral glucose (19,28). In addition, when an intravenous glucose tolerance test was performed, the results showed -866A allele carriers have a lower insulin secretion, and a reduced risk of diabetic neuropathy (29). Carriers of the -866A allele of the *UCP2* gene are reported to have a reduced risk of coronary artery disease (30), depleted energy levels in the peripheral nerve function (31), a higher waist-to-hip ratio, a high risk of metabolic syndrome (32) and higher plasma markers of oxidative stress (33).

The -866G allele was shown to be associated with a reduced mRNA expression in adipose tissue, decreased transcriptional activity, high BMI, fat mass changes (34), increased risk of obesity (17,18,21), high insulin response to glucose and a reduced risk of T2D (10,17-19). Carriers of the -866G allele of *UCP2* are reported to have low blood triglyceride levels, high insulin sensitivity (10,19) and higher levels of low-density lipoprotein (LDL) cholesterol (35). Taken together, the results showed that the -866G allele was found to increase the risk of T2D and obesity compared to the -866G/A and -866A alleles (35). Additionally, 45 insertion/deletion polymorphisms of the *UCP2* gene have been reported to be strongly associated with BMI in a Korean population (36). Taking all of the data into consideration, it is assumed that since the *UCP2* gene is associated with T2D and obesity, it is also associated with obesity in females with PCOS.

The aim of the present study was to analyze the association between the -866G/A polymorphism of the *UCP2* gene and PCOS in Korean females. In conclusion, these results indicate that no significant association exists between the -866G/A polymorphism in the promoter of the *UCP2* gene and PCOS in Korean females. The -866G/A polymorphism of *UCP2* was reported to be associated with obesity, insulin resistance and hyperinsulinemia but not associated with PCOS in patients who share common characteristic symptoms. Thus, this genetic association study provides no evidence for the involvement of the -866G/A polymorphism in the promoter of the *UCP2* gene in PCOS patients in the Korean population. Further investigation of associations between the -866G/A polymorphism of the *UCP2* gene and PCOS patients of different ethnic groups is required.

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