Lack of association between IL10 and TNFα gene polymorphisms and polycystic ovary syndrome in Saudi women

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Genet. Mol. Res. 18 (3): gmr18270
Received February 25, 2018
Accepted July 01, 2019
Published July 04, 2019
DOI http://dx.doi.org/10.4238/gmr18270

ABSTRACT. Polycystic ovarian syndrome (PCOS) is the most prevalent endocrine disorder affecting females. It is a common cause of menstrual irregularities and infertility during a woman’s reproductive years. Many factors may play a crucial role in the pathogenesis of PCOS. One of these factors is related to immunogenetics. Cytokines are significant immunomodulatory proteins for regulating and controlling cell functions involved in the immune system. The imbalance between pro- and anti-inflammatory pathways plays a role in PCOS etiology. We investigated the relationship between gene polymorphisms interleukin (IL-10) (rs1800871, rs1800872, rs1800896) and tumor necrosis (TNF-α) (rs1799724) in Saudi women with PCOS. The study group consisted of 93 Saudi females (mean age 31.05 ± 0.590, range 18 - 40 years) with PCOS. The control group consisted of 98 Saudi women without PCOS. Blood was obtained, and DNA was extracted for each patient and control. Single nucleotide polymorphisms (SNPs) of promoter regions were determined using TaqMan genotyping assays. The
polymorphism frequencies in IL-10 (rs1800871) A/A, A/G, and G/G genotypes were 7.5%, 38.7%, and 53.76%, respectively in PCOS patients and 5.1%, 45.9%, and 49.0%, respectively in controls with no significant differences between the groups. No significant differences in IL-10 polymorphism frequencies in C/C, C/T, and T/T between groups were noted. No significant differences were seen in parameters for TNF-α between groups with respect to the same TNF-α. The differences in frequencies of alleles and genotypes were not significant between Saudi women with PCOS and controls. We conclude that though in some populations, meta-analyses showed an association between IL-10 and TNF-α gene polymorphisms and PCOS, this association is not apparent in Saudi females.

Key words: IL-10; TNF-α gene polymorphism; Saudi Arabia; Polycystic ovary syndrome (PCOS)

INTRODUCTION

The most unique and habitually encountered endocrine malfunction in women is polycystic ovary syndrome (PCOS) (Qin et al., 2015; Sóter et al., 2015; Xing et al., 2017). Overproduction of androgens, ovulatory malfunctions, and other anomalies in ovarian morphology (Zhang et al., 2015) are some of the characteristics of PCOS, which is found in approximately 9–21% women of reproductive age (Yamada et al., 2017; Zeng et al., 2018). PCOS is thought of as a biological issue that impacts including other body systems, for example, those associated with psychological structure and metabolic, and procreative systems (Talaat et al., 2016). Androgen instability as the cause of syndromes involving the hypothalamus, pituitary glands, and ovaries is one of the causes of PCOS. Elevated gonadotropin activity, leading to increased levels of luteinizing hormone (LH), which stimulates ovaries to release a greater amount of androgens, mostly testosterone, is the main cause of PCOS (Szczuko et al., 2016).

PCOS is a common disease that is ascribed to a handful of hereditary and environmental risk factors. Different PCOS phenotypes may result from the interaction between a number of predisposing genomic mutations, each exerting only minor functions, and strong environmental factors (Zhang et al., 2015). PCOS is indicate by important metabolic abnormalities that include deformities of energy use, which causes obesity (Ob), glucose-induced hyperinsulinemia, and fasting, peripheral insulin resistance that disturbs the muscle and adipose tissue metabolism and also causes dyslipidemia (Wang et al., 2017; Zhang et al., 2018). It is frequently represented by hyperandrogenism with great risks of insulin resistance and abdominal obesity. It is also related to infertility, acne, hirsutism, infertility, increased risk of cardiovascular diseases, hypertension, high blood pressure, dyslipidemia, inflammation, and type 2 diabetes (Fulghesu et al., 2011; Lindholm et al., 2011; Qin et al., 2015; Wu et al., 2015; Talaat et al., 2016).

Many environmental issues, such as drinking alcohol, consuming food kept plastic packages, adrenal dysfunction, and obesity are involved in PCOS development (Sóter et al., 2015; Xing et al., 2017). The etiology of PCOS includes epigenetic changes and numerous genetic changes that are regarded as unclear; nonetheless, polymorphisms found in cytokine
IL10 and TNFα polymorphisms unassociated with PCOS in Saudis

Genes may have a significant role, particularly the possibilities for functional SNPs affecting PCOS vulnerability. Various molecular epidemiological studies have emphasized the connection of PCOS risk and cytokine (tumor necrosis factor [TNF]-α, interleukins [IL]-1A, -1B, -6, -10, and -18) gene polymorphisms (Talaat et al., 2016). This study proposes that PCO-related environmental and/or epigenetic factors interact with cytokine genes (Sóter et al., 2015). In PCOS pathogenesis, immune dysregulation could play a major role (Qin et al., 2015; Talaat et al., 2016; Xing et al., 2017). Cytokines are cell-signaling protein molecules secreted by immune cells and are engaged in many intercellular communication pathways. Discrepancies comparing pro-inflammatory and anti-bacterial cytokines and chronic infections are also believed to contribute to the history and origin of PCOS (Talaat et al., 2016).

The antenatal cytokines, which include IL-10 and -1 receptor antagonists (IL1-RA), vary with PCOS-associated inflammation (Wu et al., 2015). Regulation of inflammatory syndrome results in recruitment of regulatory B cells after excretion of IL-10 (Liu et al., 2016). IL-10 is an antenatal cytokine; in addition, it suppresses immune responses. It is also involved in the regulation of immune body processes. IL-10 was originally identified in T-helper (Th)2 cells that repress cytokine fusion in Th1 cells. IL-10 is an anti-inflammatory cytokine, during infection, it inhibits the activity of Th1 and natural killer (NK) cells and macrophages, all of which are required for optimal pathogen clearance but also contribute to tissue damage (Couper et al., 2008). PCOS patients have hyperinsulinemia and insulin-resistance, which seems to indicate an infective role for the illness. IL-10 gene polymorphism may add to PCOS pathogenesis (Karadeniz et al., 2008).

Studies on reproductive biology have revealed that these pro-inflammatory cytokines influence ovarian function in addition to the processes of fertilization, implantation, and ovulation in women with PCOS (Wu et al., 2015). In contrast to the entire menstrual cycle, the corpus luteum secretes immunoreactive TNF-alpha (Guo et al., 2015). Hyperexpression of adipose tissues in addition to TNFα in muscles are clearly programmed for an increase in insulin resistance (IR) through reduction of tyrosine kinase activity at the site of cellular insulin production. TNFα causes hyperandrogenism (HA), promotes IR, and is also associated with follicular formation; therefore, TNFα is implicated in PCOS-associated functional changes (Thathapudi et al., 2014).

Numerous studies have shown that polymorphisms in cytokine genes are associated with PCOS. Talaat et al. (2016) conducted a study in 61 patients with PCOS and 80 healthy controls in an Egyptian population and examined the influence of IL-10 serum levels and genetic polymorphisms with respect to the risk of PCOS. They found that -819 TT and IL-10 - 1082 GG genotype could be observed as a risk factors for PCOS, and IL-10 levels were meaningfully higher in standard controls compared to PCOS patients. Xing et al. (2017) reported that they observed a significant relationship between the IL-10 rs1800871 polymorphism and development of PCOS in the Chinese population.

Thathapudi et al. (2014) indicated a role for TNF-α in the pathological process of Ob in addition to IR within PCOS, and noted C850T (rs1799724) distinct phenotypes in the promoter area of the TNFα gene in a group 204 age-matched healthy controls and 204 PCOS patients. This study’s outcome suggests that the TNFα system results in initiation of IR, HA, and Ob in PCOS patients irrespective of the discrepancies of the TNFα C850T (rs1799724) in the Indian women. Sóter et al. (2015) performed a study in 196 age-matched women with 97 healthy women as controls and 99 with PCOS. Using polymerase chain
reaction (PCR), the authors were able to show a correlation between IL-10 and -6, interferon (IFN)y, and transforming growth factor (TGF)b1 distinct phenotypes associated with genes encoding inflammation-associated factors from peripheral blood-derived DNA and cytokines. They noticed that distinct gene cytokine-associated phenotypes were not found in Brazilian women with PCOS development; nevertheless, these phenotypes may result in common metabolic complaints associated with PCOS.

The association of the polymorphisms of the TNFα and IL-10 and -6 genes with PCOS occurrence and the clinical/laboratory characteristics in Swedish residents was described in detail. Vural et al. (2010) designed a study with SNPs of the TNF-α (-308 G/A), IL-6 (-174 G/C), IL-10(-1082 G/A) genes in DNA 95 healthy controls and in peripheral white blood cells of 97 patients with PCOS. They showed that there was a trend toward a higher rate of normal genotype in the controls compared to the C allele and IL-6 CC genotype among PCOS women although this difference was not expected. There were no or very small difference observed between groups in allele or genotype frequencies for TNFα and IL-10 genes. Yun et al. (2011) performed a study with 144 healthy women as controls out of a total of 217 study patients. They did a comparative study of the -1031(T/C) genetic variation in the TNFα gene with PCOS in Korean women; they found a relationship in PCOS with the -1031(T/C) polymorphism in the promoter region of the TNF α gene (P-value = 0.0003; odds ratio [OR] = 2.53). However, controls were smaller in number than the C allele PCOS patients.

Karadeniz et al. (2008) studied the IL-10 polymorphism in a study of Turkish women, consisting 74 healthy controls and 91 young women with PCOS. They found that the IL-10 gene polymorphism of PCOS patients was not involved with changes in inflammatory markers.

In our study, we examined the relationship between the gene polymorphisms IL-10 (rs1800871, rs1800872, rs1800896) and TNFα (rs1799724) in Saudi women with PCOS.

MATERIAL AND METHODS

Subjects

This study included 93 Saudi women (mean age 31.05 ± 0.59 years) who had PCOS and had presented to the King Khaled University Hospital, Riyadh, Saudi Arabia. The control group included 98 (mean age 31.48 ± 0.49 years) subjects. Ethical approval for the study was obtained from the Medical Ethics Committee of King Khalid University Hospital and the Ethical Committee of King Saud University. All patients and controls provided informed consent and agreed to give blood samples for this case-control study.

DNA preparation

The study group consisted of 93 Saudi females (mean age 31.05 ± 0.590, range 18 - 40 years) with PCOS, and the controls consisted of 98 non-PCOS Saudi women. Blood was obtained and genomic DNA was extracted from peripheral blood using the Puregene purification kit (Qiagen; Hilden, Germany) according to the manufacturer’s protocol. Quantification of extracted DNA was performed using a NanoDrop ND-2000c spectrophotometer (Thermo Scientific, Wilmington, DE, USA).
Genotyping of IL-10 rs1800871, rs1800872, rs1800896, and TNF-a rs1799724

Subjects were genotyped for polymorphism IL-10 rs1800871, rs1800872, rs1800896, and TNF-a rs1799724 using the TaqMan® SNP Genotyping Assay (Applied Biosystems Inc., Foster City, CA, USA) on an ABI 7500 real-time PCR (qPCR) System (Applied Biosystems) for detection of IL-10 rs1800871, rs1800872, rs1800896, and TNFα rs1799724. A 25 μL PCR reaction mixture consisted of 1X TaqMan® Genotyping Master Mix (Applied Biosystems), 1X SNP Genotyping Assay Mix, and 50 ng DNA. Each 96-well plate included two no-template controls. qPCR was performed on an ABI 7500 system using the recommended conditions consisting of incubation at 95°C for 10 min followed by 40 cycles, denaturation at 92°C for 15 s, and annealing/extension at 60°C for 1 min. The VIC® and 6-carboxy-fluorescein (FAM) fluorescence levels of the PCR products were measured at 60°C for 1 min. Analysis of fluorescence using the automated 2-color allele discrimination software on ABI 7500 showed clear discrimination of the two genotypes on a two-dimensional graph.

Statistical analysis

PCOS prevalence was estimated using sample proportion in order to see the relationship between the frequency in case and non-case groups, and for gene polymorphisms, the crude risk ratio (RR) and the crude odds ratio (OR) were used. A chi-squared test for dependency ($\chi^2$) used to assess the frequencies of IL-10 and TNFα gene polymorphisms. Logistic regression was used to examine the effects of gene polymorphisms on PCOS. A replicated stratification analysis was utilized in order to estimate the adjusted RR and adjusted OR to ensure that there was no confounders for age and body mass index (BMI). All statistical analyses were conducted using IBM-SPSS (version 22).

RESULTS

Demographic characteristics of patients and controls

Clinical characteristics and statistics of PCOS patients and controls participating in this study are outlined in Table 1. The PCOS and the control groups had no significant differences in age.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Control Mean ± SEM</th>
<th>PCOS Mean ± SEM</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (yrs)</td>
<td>31.48±0.489</td>
<td>31.05±0.590</td>
<td>0.572</td>
</tr>
<tr>
<td>Height (cm)</td>
<td>158.35±0.628</td>
<td>158.16±0.539</td>
<td>0.817</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>71.57±1.37</td>
<td>69.25±1.28</td>
<td>0.213</td>
</tr>
<tr>
<td>BMI (kg/m2)</td>
<td>28.54±0.522</td>
<td>27.55±0.528</td>
<td>0.290</td>
</tr>
</tbody>
</table>

SEM= Standard Error of Mean; P=Significance
IL-10 gene (rs1800871) A/G polymorphism

The prevalence of IL-10 polymorphisms was determined for Saudi women (98 PCOS patients and 93 controls). Table 2 presents the base pairs in the wild type homozygous (AA), heterozygous (AG), and mutated homozygous (GG). Overall, the distribution of the various genotypes of IL-10 (rs1800871) did not differ significantly between PCOS patients and controls (P = 0.796).

<table>
<thead>
<tr>
<th>Genotype</th>
<th>Control No.</th>
<th>PCOS No.</th>
<th>Control vs. Patients</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>OR</td>
</tr>
<tr>
<td>AA</td>
<td>5</td>
<td>7</td>
<td>0.661</td>
</tr>
<tr>
<td>AG</td>
<td>45</td>
<td>36</td>
<td>0.571</td>
</tr>
<tr>
<td>GG</td>
<td>48</td>
<td>50</td>
<td>0.826</td>
</tr>
<tr>
<td>TOTAL</td>
<td>98</td>
<td>93</td>
<td>0.677-1.663</td>
</tr>
</tbody>
</table>

Table2. IL-10 (rs1800871) A/G genotype in 98 polycystic ovary syndrome (PCOS) patients compared to 93 controls.

IL-10 gene (rs1800872) T/G polymorphism

The prevalence of the IL-10 polymorphism was determined for Saudi women (98 PCOS patients and 93 controls). Table 3 presents the base pairs in mutated homozygous (GG), heterozygous (TG), and also the wild type homozygous (TT). The overall, distribution of the various genotypes of IL-10 (rs1800872) did not differ significantly between PCOS and controls (P = 0.693)

<table>
<thead>
<tr>
<th>Genotype</th>
<th>Control No.</th>
<th>PCOS No.</th>
<th>Control vs. PCOS</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>OR</td>
</tr>
<tr>
<td>TT</td>
<td>5</td>
<td>5</td>
<td>0.946</td>
</tr>
<tr>
<td>TG</td>
<td>43</td>
<td>37</td>
<td>0.860</td>
</tr>
<tr>
<td>GG</td>
<td>50</td>
<td>51</td>
<td>0.858</td>
</tr>
<tr>
<td>TOTAL</td>
<td>98</td>
<td>93</td>
<td>0.694-1.731</td>
</tr>
</tbody>
</table>

Table3. IL-10 (rs1800872) T/G genotype in polycystic ovary syndrome (PCOS) patients compared to controls.

IL-10 gene (rs1800896) T/C polymorphism

No.: Number of individuals, OR: Odds Ratio, CI : Confidence Interval, \(\chi^2\): Chi Square
The prevalence of IL-10 polymorphism was determined for Saudi women (98 PCOS patients and 93 controls). Table 4 presents the frequencies of the wild type homozygous (TT), heterozygous (TC), and mutated homozygous (CC). Overall, the distribution of the various genotypes of IL-10 (rs1800896) did not differ significantly between PCOS patients and controls (P = 0.141).

<table>
<thead>
<tr>
<th>Genotype</th>
<th>Control No.</th>
<th>PCOS No.</th>
<th>Control vs. PCOS OR</th>
<th>CI</th>
<th>( \chi^2 )</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>TT</td>
<td>42</td>
<td>28</td>
<td>1.741</td>
<td>0.958-3.163</td>
<td>0.13</td>
<td>0.953</td>
</tr>
<tr>
<td>TC</td>
<td>38</td>
<td>46</td>
<td>1.816</td>
<td>0.955-3.453</td>
<td>3.33</td>
<td>0.067</td>
</tr>
<tr>
<td>CC</td>
<td>18</td>
<td>19</td>
<td>0.876</td>
<td>0.427-1.797</td>
<td>0.13</td>
<td>0.718</td>
</tr>
<tr>
<td>TOTAL</td>
<td>98</td>
<td>93</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Table 4. IL-10 (rs1800896) T/C genotype in polycystic ovary syndrome (PCOS) patients compared to controls.**

No.: Number of individuals, OR: Odds Ratio, CI: Confidence Interval, \( \chi^2 \): Chi Square

**TNF-a gene (rs1799724) C/T polymorphism**

The prevalence of the IL-10 polymorphism was determined for Saudi women (98 PCOS patients and 93 controls). Table 5 presents the base pairs in the wild type homozygous (CC), heterozygous (CT), and mutated homozygous (TT). Overall, the distribution of the various genotypes of IL-10 (rs1799724) did not differ significantly between PCOS and control groups (P = 0.953).

<table>
<thead>
<tr>
<th>Genotype</th>
<th>Control No.</th>
<th>PCOS No.</th>
<th>Control vs. PCOS OR</th>
<th>CI</th>
<th>( \chi^2 )</th>
<th>P - value</th>
</tr>
</thead>
<tbody>
<tr>
<td>CC</td>
<td>85</td>
<td>80</td>
<td>0.992</td>
<td>0.992-1.000</td>
<td>0.02</td>
<td>0.885</td>
</tr>
<tr>
<td>CT</td>
<td>12</td>
<td>13</td>
<td>1.062</td>
<td>0.465-2.430</td>
<td>0.02</td>
<td>0.885</td>
</tr>
<tr>
<td>TT</td>
<td>1</td>
<td>0</td>
<td>1.151</td>
<td>0.496-2.671</td>
<td>0.02</td>
<td>0.885</td>
</tr>
<tr>
<td>TOTAL</td>
<td>98</td>
<td>93</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Table 5. TNF-a (rs1799724) C/T genotype in polycystic ovary syndrome (PCOS) compared to controls.**

No.: Number of individuals, OR: Odds Ratio, CI: Confidence Interval, \( \chi^2 \): Chi Square

**DISCUSSION**

Lymphocytes and monocytes produce IL-10 that has been shown to be an essential anti-inflammatory compound and immune-balancing cytokine among others because it efficiently down-regulates pro-inflammatory cytokines, such as IL-1 and -6 and TNFα. Since an imbalance in pro- and anti-inflammatory cytokines could participate in PCOS, the
polymorphic sequence of IL-10 may be a crucial biomarker for PCOS predisposition (Karadeniz et al., 2008; Talaat et al., 2016).

Our results were compared with those reported in literature and showed that our findings were in agreement with those of several studies that indicated that numerical combinations of genetic polymorphism of IL-10 do not influence PCOS (Karadeniz et al., 2008; Vural et al., 2010; Lindholm et al., 2011; Sóter et al., 2015). Results by Talaat et al. (2016) did not agree with our findings in a study in 80 controls in relation to 61 patients with PCOS in Egyptian women. They examined the influence of genetic polymorphisms in addition to IL-10 serum levels as risks for PCOS. Results showed that IL-10 -1082 GG and -819 TT genotype could be considered one of the causes for PCOS, and IL-10 levels in normal controls were higher compared to PCOS patients. A study by Xing et al. (2017) reported an association between growth of PCOS in the Chinese population and the IL-10 rs1800871 polymorphism.

Numerous ailments are regulated by TNFα, which is a multifunctional pro-inflammatory cytokine. It exists not only in oocytes and granulose cells, but also in human ovarian follicular fluid. TNFα is believed to be connected with anovulation, increased ovarian steroid secretion, and ovarian apoptosis (Yun et al., 2011).

The results for the TNFα (rs1799724) C/T polymorphism also showed that the majority of female patients and controls were homozygous for CC, and about 12% of the controls and 14.0% of PCOS patients existed in the heterozygous (CT) state and also in the wild type allele (CC). Nevertheless, PCOS and control groups did not significantly different.

Comparison of our results with those reported in literature indicated that, our analysis agreed with one study in which it was inferred that numerical combinations of genetic polymorphisms of TNFα had no relationship with PCOS (Vural et al., 2010; Gou et al., 2015). On the other hand, there is a contrasting study by Yun et al. (2011) in which 217 PCOS patients and 144 matched female controls (healthy women) were studied. A comparative study of the -1031(T/C) polymorphism of the TNFα gene with PCOS in a Korean population was done, and the outcome showed that there was a strong relationship between PCOS (P - value = 0.0003, OR = 2.53) and the -1031(T/C) polymorphism in the promoter region of the TNFα gene. Moreover, the occurrence of the C allele was lower in controls in comparison to PCOS patients. Thathapudi et al. (2014) reported that the TNFα system might contribute to the pathogenesis of HA, Ob, and IR in PCOS patients independent of the polymorphism of the TNF-α C850T (rs1799724) as shown in an Indian population.

However, our results do not mean that a relationship between SNPs and PCOS does not exist; differences may be due to differences in life styles. There are no reports on the genetic polymorphisms in cytokine genes in Saudi PCOS patients, and we need more research concerning cytokines associated with PCOS, especially in the Saudi population whose women are the most vulnerable.

ACKNOWLEDGMENTS

The authors would like to thank the Deanship of Scientific Research at King Saud University for funding this study through the Research Project.
CONFLICTS OF INTEREST

The authors declare no conflict of interest.

REFERENCES


