IL-1α C376A Transversion Variant and Risk of Idiopathic Male Infertility in Iranian Men: A Genetic Association Study

Tayyebeh Zamani-Badi, M.Sc.1, Mohammad Karimian, Ph.D.2, Abolfazl Azami Tameh, Ph.D.2, Hossein Nikzad, Ph.D.2*
1. Gametogenesis Research Center, Kashan University of Medical Sciences, Kashan, Iran
2. Anatomical Sciences Research Center, Kashan University of Medical Sciences, Kashan, Iran

Abstract

Background: IL-1α produced by Sertoli cells is considered to act as a growth factor for spermatogonia. In this study, we investigated the association of the C376A polymorphism in IL-1α with male infertility in men referring to the Kashan IVF Center.

Materials and Methods: In this case-control study, 2 ml of blood was collected from 230 fertile and 230 infertile men. After DNA extraction, the C376A variant was genotyped by polymerase chain reaction-restriction fragment length polymorphism (PCR-RFLP). In addition, the molecular effects of the C376A transversion were analysed using bioinformatics tools.

Results: A significant association was observed between the homozygous genotype CC with male infertility [odds ratio (OR)=1.97, 95% confidence interval (CI)=1.14-3.41, P=0.016]. Carriers of C (AC+CC) showed a similar risk for male infertility (OR=1.78, 95% CI=1.06-2.99, P=0.030). Also, allelic analysis showed that the C allele is associated with male infertility (OR=1.43, 95% CI=1.09-1.88, P=0.011). In sub-group analysis, we found that the AC genotype is associated with asthenozoospermia (OR=2.38, 95% CI=1.03-5.53, P=0.043). In addition, carriers of C were at high risk for asthenozoospermia (OR=2.25, 95% CI=1.03-4.10, P=0.047). Also, C allele was significantly associated with oligozoospermia (OR=1.44, 95% CI=1.01-2.06, P=0.049) and non-obstructive azoospermia (OR=1.67, 95% CI=1.04-2.68, P=0.034). Finally, in silico analysis showed that the C376A polymorphism could alter splicing especially in the acceptor site.

Conclusion: This is the preliminary report on the association of IL-1α C376A polymorphism with male infertility in the Kashan population. This association shows that the IL-1α gene may be a biomarker for male infertility, and therefore needs additional investigations in future studies to validate this.

Keywords: Genetic Polymorphism, Interleukin-1α, Male Infertility, Spermatogenesis

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Introduction

Male infertility is a multifactorial syndrome that affects up to 12% of men (1). Male factors are responsible for 40-50% of total infertility cases (2). In more than 70% of cases, there is a conclusive reason including varicocele, aneuploidies, infectious diseases and post-testicular obstruction, however, in less than 30% of infertile males, the cause of their infertility is unknown and are thus diagnosed as idiopathic (3, 4).

Environmental, lifestyle, physiological and genetic factors are involved in male infertility (5-7). From numerous genetic factors that are essential for normal spermatogenesis, cytokines play an important role (8). These are regulatory peptides which regulate testicular and glandular function (9).

Human seminal plasma contains several cytokines including IL-1, IL-2, IL-4, IL-5, IL-6, IL-7, IL-10, IL-11, IL-12, IL-13, IL-17, IL-18, IL-23, TNFα, IFN-γ, TGFα, TGFβ (8). One of the most important gene sets involved in fertility is the interleukin-1 (IL-1) gene family which encodes regulatory cytokines playing multifaceted roles in the male reproductive system. For example, they may act as growth factors and are involved in physiological protection, germ cell proliferation and differentiation, regulation of junctions and steroidogenesis (10, 11).

The IL-1 gene family members include IL-1α (OMIM: 147760), IL-1β (OMIM: 147720) and IL-1RA (OMIM: 147679), all of which are located on chromosome 2q14 (12). IL-1α is secreted from seminiferous epithelium and is known as a growth factor for immature Sertoli cells and...
spermatogonia (13).

Single nucleotide polymorphisms (SNPs), by altering the structure of genes involved in spermatogenesis, may affect gene expression, mRNA structure and protein function, and may therefore lead to male infertility (14-16). Therefore, evaluating SNPs in the IL-1 gene family could be considered as an interesting research topic. A SNP (C376A; rs2071376) has been found to have a high frequency in the IL-1α gene. The association of this SNP with some disorders has been investigated in different studies including cancers (17, 18), systemic sclerosis (19), periodontitis (20), endometriosis (21) and keraotoncus (22). The association between the C376A SNP and idiopathic male infertility has, however, not been reported. In this study, we investigated the association between the IL-1α C376A SNP and idiopathic male infertility in an Iranian population as a preliminary project. Also, we evaluated the functional effects of C376A on IL-1α using bioinformatics tools.

Materials and Methods

Subjects and inclusion criteria

In this cross-sectional study, a total of 460 samples comprising 230 infertile men (with mean age of 30.93 ± 5.47) and 230 fertile men (with mean age of 32.12 ± 5.52) selected among individuals attending the Kashan Infertility Centre (Shahid Beheshti Hospital, Kashan, Iran). Infertile patients were defined as "idiopathic" and selected based on andrological examination. Patients with previous testis trauma, obstruction of the vas deferens, infectious and chronic diseases, hypogonadotropic hypogonadism, abnormal hormonal profile (Luteinizing, Follicle Stimulating, and testosterone hormones) and abnormal karyotype or Y chromosome microdeletions were excluded from the study. According to the World Health Organization (WHO) 1999 criteria, the patient sub-groups were determined (23) and the subjects were categorized into non-obstructive azoospermia (n=51) without spermatozoa in the ejaculated semen, oligozoospermia (n=95) with sperm concentration less than 20 million/ml, and asthenozoospermia (n=84) with progressive sperm motility less than 50%.

The control group was randomly selected from healthy men referred to the Kashan Infertility Centre. They had normal sperm parameters, had no history of chronic and familial diseases and had at least one offspring. Finally, a total of 2 ml of whole blood was collected from all males into EDTA-K3 containing tubes and were stored in -20°C for further usage. Written informed consent was obtained from all case and control subjects. The study was approved by the Medical Research Ethics Committee of the Kashan University of Medical Sciences (IR.KAUMS.REC.1394.6).

Single nucleotide polymorphism genotyping

Total genomic DNA was isolated from whole blood by using a DNA extraction kit (Bioneer, Korea). Purified DNA was stored at -20°C for further use. The IL-1α C376A SNP was genotyped by the polymerase chain reaction-restriction fragment length polymorphism (PCR-RFLP) method. For this purpose, forward and reverse primers flanking the SNP were designed based on the complete sequence of IL-1α by the Oligo7 software (Molecular Biology Insights, Inc., Cascade, CO, USA). The sequences of the primers were: 5’-ATGCTAAATATTACCGTGATTCT-3’ 5’-AGATCAATGGAGAAATGCGATT-3’ respectively.

The PCR was carried out in a total volume of 20 µl containing 10µl pre-mix (CinnaGen, Iran), 0.35 µM of each forward and reverse primers, and 3 µl of template DNA. PCR cycling conditions were an initial denaturation step at 94°C for 5 minutes followed by 35 cycles of denaturation at 94°C for 45 seconds, annealing at 56.9°C for 1 minute and extension at 72°C for 1 minute along with a final extension at 72°C for 5 minutes. PCR products were then digested with the BstYI restriction enzyme (CinnaGen, Iran). For this purpose, approximately 0.1 µg of the PCR product was incubated with 5 units of BstYI at 37°C for 16 hours. Finally, BstYI was inactivated by incubation at 65°C for 20 minutes. The digested fragments were separated on a 1% agarose gel stained with DNA Green Viewer (CinnaGen, Iran) and visualised under the UV light. To verify PCR-RFLP results, 2% of samples were sequenced randomly. PCR product recovery kit (Roche Applied Science, Mannheim, Germany) was used to purify the PCR product (368 bp in length). Direct sequencing of the purified PCR products was undertaken by Bioneer (Daejeon, Korea). Chromas (version 2.33) was used to check the chromatograms.

Statistical analysis

The difference in frequencies of genotypes and alleles between the case and control groups was analyzed by Chi-square test. For association analysis, the odds ratios (ORs) and 95% confidence intervals (95% CI) were estimated by a binary regression logistic test. A two-tailed p-value less than 0.05 (P<0.05) was considered significant. All analyses were conducted in the SPSS software (SSPS Inc., IBM Corp, Armonk, NY, USA) version 19.

In silico analysis

Bioinformatics tools were used to analyze the influence of the IL-1α C376A intronic SNP on RNA structure and splicing pattern. The effect on RNA structure and splicing was assessed with RNAspn online server (24) and NetGene2 (25) respectively. Finally, reported interactions of IL-1α with other molecules were obtained from the BioGRID interactome database (26).

Results

Polymerase chain reaction-restriction fragment length polymorphism and DNA sequencing

Results of PCR-RFLP showed that 368 bp fragment was fully digested into 114 bp and 254 bp fragments in some
samples, showing the efficiency of the method used. The samples with two, three and one fragments were identified as CC, AC, and AA genotypes respectively (Fig.1A). The data from direct sequencing also confirmed the results of PCR-RFLP (Fig.1B).

**IL-1α C376A distribution**

In this study, the genotype and allele frequencies of the IL-1α C376A SNP were compared between the infertile and healthy groups (Table 1). We observed a significant association between the homozygous genotype CC with male infertility (OR=1.97, 95% CI=1.14-3.41, P=0.016). Carriers of C (AC+CC) were at a similar risk for male infertility (OR=1.78, 95% CI=1.06-2.99, P=0.030). Also, allelic analysis showed that the C allele is associated with infertility (OR=1.43, 95% CI=1.09-1.88, P=0.011). In sub-group analysis, we found that the AC genotype is associated with asthenozoospermia (OR=2.38, 95% CI=1.03-5.53, P=0.043). In addition, there was a significant association between carriers of C and asthenozoospermia (OR=2.25, 95% CI=1.01-4.10, P=0.047). Also, C allele was significantly associated with oligozoospermia (OR=1.44, 95% CI=1.01-2.06, P=0.049) and non-obstructive azoospermia (OR=1.67, 95% CI=1.04-2.68, P=0.034).

**In silico analysis**

Functional consequence of the C376A transversion on RNA structure was evaluated. However, no significant effect on RN (distance: 0.0191, P=0.686) was observed (Fig.2). Minimum free energy of normal RNA was equal to -81.80 kcal/mol but increased to -80.50 kcal/mol for the variant allele. The data from NetGene2 revealed that the C370A SNP alters the IL-1α splice site pattern on the direct strand (+ strand) especially for the acceptor splice pattern (Fig.2). The BioGRID interactome showed that IL-1α has 17 gene-gene interactions (Fig.3).
In this study, we examined the association of the IL-1α C376A SNP with male infertility in an Iranian population (Kashan, Iran) as a pilot study. Our study revealed that not only the CC genotype was associated with male infertility, but also the C allele showed significant association. In addition, carriers of the C allele were at almost two-fold risk for male infertility. Sub-group analysis revealed that AC genotype and carriers of C were associated with asthenozoospermia. Also, the C allele was significantly associated with oligozoospermia and non-obstructive azoospermia. Therefore, IL-1α C376A is a potential genetic risk factor for male infertility, although further studies of different ethnicities in Iran and other populations are required to obtain a more accurate picture. After Hardy-Weinberg equilibrium (HWE) calculation in the control group, we found a highly significant deviation. However, the case group showed no deviation. even though it does not necessarily need to follow HWE due to the inherent sampling bias in cases. The deviation from HWE in the control group (normozoospermic men) could also be due to the selection bias (27) given that not all men in the general population will be fertile.

Spermatogenesis is a dynamic process in which many factors are necessary for creating and regulating balance in this process. For example, growth factors and cytokines are essential for development of functional spermatozoa (28, 29). Interleukin-1 is produced by epithelia of seminiferous tubules and acts as a physiological paracrine/autocrine factor on testicular cells and required for immunological protection (30). There is a probable mechanism that in the absence of testosterone, followed by increased cell apoptosis, spermatogenesis is finally reduced (31, 32). The second probable mechanism is excess reactive oxygen species (ROS).

The presence of the associated SNP and the consequent change in the amount of interleukin along with excess production of ROS may reduce sperm motility. One of the reasons for reduced sperm motility may be DNA damage and lipid peroxidation of sperm membrane (33). Also, increased ROS with oxidizing DNA or proteins, enzyme inhibition, cell death and apoptosis of sperm may cause the oligozoospermia phenotype (34, 35). Due to these possible mechanisms, the association of the IL-1α SNP with some abnormalities in sperm parameter may be explained. SNPs could change the gene expression pattern (14), mRNA structure (36, 37), splicing pattern (38) and protein function (39, 40). In silico tools, which can predict the damaging effects of SNPs, were therefore used especially that IL-1α C376A is an intronic SNP and may affect RNA structure and splicing. Although we found no evidence for C376A to affect RNA structure, we observed a predicted effect on splicing alteration. Therefore, the association of this SNP may be due to this effect. In this study, there were various limitations including gene-environment and gene-gene interactions that must be consid-

### Table 1: Allelic and genotypic distribution of the IL-1α C376A SNP

| Genotype/Allele | Control n=230 | All cases n=282 | Oligo n=95 | Asteno n=84 | NOA n=51 | Total | OR (95% CI) | P value |
|----------------|--------------|----------------|-----------|------------|---------|-------|------------|---------|
| AA            | 44 (19.13)   | 27 (11.74)     | 14 (14.74)| 8 (9.52)   | 5 (9.80)| -     | -         | -       |
| AC            | 90 (39.13)   | 87 (37.83)     | 30 (31.58)| 39 (46.43)| 18 (35.29)| 1.58  | 1.05  (0.90-2.74) | 1.76  (0.61-5.05) | 0.113 0.901 0.043 0.293 |
| CC            | 96 (41.74)   | 116 (50.43)    | 51 (53.68)| 37 (44.05)| 28 (54.90)| 1.97  | 1.67  (1.14-3.41) | 2.12  (0.91-4.93) | 0.016 0.146 0.081 0.069 |
| AC+CC         | 186 (80.87)  | 203 (88.26)    | 81 (85.26)| 76 (90.48)| 46 (90.20)| 1.78  | 1.37  (1.06-2.99) | 2.25  (0.71-2.64) | 0.030 0.348 0.047 0.127 |
| A             | 178 (38.70)  | 141 (30.65)    | 58 (30.53)| 5 (32.74) | 28 (27.45)| -     | -         | -       |
| C             | 282 (61.30)  | 319 (69.35)    | 132 (69.47)| 113 (67.26)| 74 (72.55)| 1.43  | 1.44  (1.09-1.88) | 1.30  (0.89-1.88) | 0.011 0.049 0.172 0.034 |

SNP: Single nucleotide polymorphism, OR: Odds ratio, Oligo: Oligozoospermia, Asteno: Asthenozoospermia, and NOA: Non-obstructive azoospermia.

Significant differences between the case and control groups are shown in bold type.
eroded in subsequent studies. Also, lack of in vitro studies such as investigating the effect of the SNP on IL-1α gene expression and isoform formations due to splicing alterations is another limitation of this study.

Conclusion

Our study suggests that the IL-1α C376A SNP may increase the risk of male infertility up to two-fold. Since this is the first study, future studies with larger sample sizes in different ethnicities and populations is warranted given the variable environmental factors in different geographic regions.

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Author’s Contribution

M.K., H.N.; Planned and supervised the study upon which the current subset project was based. M.K., T.Z.-B.; Developed the outline for the current study and supervised the analysis of the samples. T.Z.-B.; Write the manuscript and M.K., H.N. revised the paper. M.K., H.N., T.Z.-B., A.A.T.; Contributed to data analysis and prepared the manuscript. All authors reviewed and approved the final manuscript.

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