Effect of angiotensin I-converting enzyme and α-actinin-3 gene polymorphisms on sport performance

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Abstract. Genetic polymorphism is considered to be associated with human physical performance. The angiotensin I-converting enzyme insertion/deletion (ACE I/D) and the α-actinin-3 (ACTN3) R577X polymorphisms have been widely investigated for such associations, and functional ACE I/D and ACTN3 R577X polymorphisms have been associated with sprinter performance. The aim of this study was to determine the effect of these polymorphisms on sport performance among 37 elite athletes and 37 healthy controls. The ACE II genotype was identified in 32.43% of the control group and 8.11% of elite athletes, the DD genotype in 37.84% of the control group and 51.35% of the elite athletes, and the ID genotype in 29.73% of the control group and 40.54% of the elite athletes. With regard to the ACTN3 gene, the XX genotype, which confers an advantage for endurance activities, was identified in 10.81% of the control group and 35.14% of the elite athletes. The XX genotype was observed more frequently than the RR genotype (advantageous for sprinting), which was identified in 2.70% of the control group and 10.81% of elite athletes. The RX genotype (observed in 86.48% of the control group and in 54.05% of the elite athletes) was the most common genotype of the individuals in the present study. The study showed that ACTN3 and ACE gene polymorphisms have an effect on muscle power; however, larger studies are required.

Introduction

Genetics is involved in determining the athletic ability of an individual. A large number of studies have revealed various associations among specific gene polymorphisms, physical ability in various sport fields, and different ethnicities and genders (1). Two genes which have been extensively investigated in association with athletic ability are angiotensin I-converting enzyme (ACE) and α-actinin-3 (ACTN3). ACE is a key component of the renin-angiotensin-aldosterone system, generating the vasoconstrictor hormone angiotensin II and degrading the vasodilator kinins (2). The ACE I/D polymorphism was the first specific gene variant to be associated with human physical performance (3). A variation in the structure of the human ACE gene (17q22-q24) has been reported in which the insertion (I) variant was associated with lower ACE levels than the deletion (D) variant (4). An excess of the I allele has been associated with certain aspects of endurance performance (5,6). The I allele is functionally associated with reduced ACE serum levels and activity. This allele is considered a favourable mutation as lower ACE activity results in less vasoconstriction and thus an increased delivery of oxygenated blood to the working muscles. Therefore, individuals that possess an I allele or the II genotype are thought to possess a greater advantage in endurance activities, including running, cycling and swimming, where the demand for oxygen is crucial. In 2009, Ahmetov et al showed that the ACE D allele was associated with greater strength and muscle volume at baseline and an increased percentage of fast-twitch muscle fibres. In addition, the D allele was shown to be associated with elite power athlete status (6).

It is generally accepted that muscle fibre composition may influence physical performance. The ACTN3 gene R577X polymorphism is suspected to be one of the contributing gene variations in the determination of muscle fibre type composition and athletic status (7). The C>T polymorphism at nucleotide position 1747 in the ACTN3 coding sequence converts an arginine (R) to a stop codon (X) at residue 577 (8). This variation generates two different versions of the ACTN3 gene common in the general population: The 577R allele is functionally associated with reduced ACE serum levels and activity. This allele is considered a favourable mutation as lower ACE activity results in less vasoconstriction and thus an increased delivery of oxygenated blood to the working muscles. Therefore, individuals that possess an I allele or the II genotype are thought to possess a greater advantage in endurance activities, including running, cycling and swimming, where the demand for oxygen is crucial. In 2009, Ahmetov et al showed that the ACE D allele was associated with greater strength and muscle volume at baseline and an increased percentage of fast-twitch muscle fibres. In addition, the D allele was shown to be associated with elite power athlete status (6).

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Key words: polymorphism, sport performance, athletic ability
The present study investigated the effect of ACE and ACTN3 gene polymorphisms in 37 elite athletes and 37 healthy controls to identify the potential correlations between genotype and physical performance.

Materials and methods

Ethical approval. The study was approved by the Ethics Committee of Istanbul University (Istanbul, Turkey; project no. 2011/2115-897).

Participants. A total of 37 healthy control individuals, aged 21.1±3.1 years, and 37 elite athletes, aged 21.1±3.1 years, were involved in the present study. The control individuals were not engaged in physical activity on a regular basis. Written informed consent was obtained from the patient/the family of the patients.

Genotyping. Molecular genetic analyses were performed using DNA samples, which were obtained from peripheral EDTA-treated blood with a DNA Extraction kit (Agilent Technologies, Inc., Santa Clara, CA, USA) according to the manufacturer’s instructions. The ACE and ACTN3 polymorphisms were amplified using polymerase chain reaction (PCR). The ACE ID polymorphism PCR was conducted using the ACE forward and reverse primers (Table I; Tib MolBiol Syntheselabor GmbH, Berlin, Germany). PCR was performed by one cycle of denaturation at 95°C for 3 min, annealing at 55°C for 50 sec, and extension at 72°C for 50 sec, and 39 cycles of denaturation at 95°C for 50 sec, annealing at 55°C for 50 sec and extension at 72°C for 50 sec, followed by a final extension step of 2 min at 72°C. ACTN3 exon 16 was amplified by PCR using ACTN3 forward and reverse primers (Table I; Tib MolBiol Syntheselabor GmbH). PCR was performed at 95°C for 5 min, 35 cycles of denaturation at 95°C for 30 sec, annealing at 60°C for 30 sec and extension at 72°C for 30 sec, and a final extension step for 10 min at 72°C. The resulting ACTN3 gene PCR products were genotyped (in the Human Diseases and Diagnosis Laboratory, Istanbul University, Istanbul, Turkey) by restriction fragment length polymorphism using Ddel restriction enzyme (Tib MolBiol Syntheselabor GmbH). Products were separated by a bioanalyser (Agilent 1000; Agilent Technologies, Inc.).

Results

Bioanalyser results. Each genotype of the ACTN3 gene was characterised using the bioanalyser. The XX genotype (Fig. 1) exhibited three peaks (86, 97 and 108 bp), the RX genotype (Fig. 2) four peaks (86, 97, 108 and 205 bp) and the RR genotype (Fig. 3) two peaks (86 and 205 bp). With regard to the ACE gene, the DD genotype exhibited one peak at 180 bp.
(Fig. 4), the II genotype one peak at 490 bp (Fig. 5) and the ID genotype two peaks at 180 and 490 bp (Fig. 6).

**Genotype frequencies.** The ACE I/D genotype frequencies in the elite athletes and the control group are shown in Table II and Fig. 7. The frequencies of the ACTN3 R577X genotypes

<table>
<thead>
<tr>
<th>Genotype</th>
<th>Controls (37)</th>
<th>Elites (37)</th>
</tr>
</thead>
<tbody>
<tr>
<td>DD</td>
<td>14 (37.84)</td>
<td>19 (51.35)</td>
</tr>
<tr>
<td>ID</td>
<td>11 (29.73)</td>
<td>15 (40.54)</td>
</tr>
<tr>
<td>II</td>
<td>12 (32.43)</td>
<td>3 (8.11)</td>
</tr>
</tbody>
</table>

ACE: angiotensin I-converting enzyme.
Table IV. Number and frequency (%) of ACTN3 and ACE genotypes in the elite athletes and controls.

<table>
<thead>
<tr>
<th>Genotype (ACTN3/ACE)</th>
<th>Group (n)</th>
<th>RR/II</th>
<th>RR/ID</th>
<th>RR/DD</th>
<th>RX/II</th>
<th>RX/ID</th>
<th>RX/DD</th>
<th>XX/II</th>
<th>XX/ID</th>
<th>XX/DD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Controls (37)</td>
<td>0</td>
<td>1 (2.70)</td>
<td>0</td>
<td>11 (29.73)</td>
<td>8 (21.62)</td>
<td>13 (35.14)</td>
<td>1 (2.70)</td>
<td>2 (5.41)</td>
<td>1 (2.70)</td>
<td></td>
</tr>
<tr>
<td>Elites (37)</td>
<td>0</td>
<td>1 (2.70)</td>
<td>3 (8.11)</td>
<td>2 (5.41)</td>
<td>9 (24.32)</td>
<td>9 (24.32)</td>
<td>1 (2.70)</td>
<td>5 (13.51)</td>
<td>7 (18.92)</td>
<td></td>
</tr>
</tbody>
</table>

ACTN3, α-actinin-3; ACE, angiotensin I-converting enzyme.

Discussion

Sport performance is likely to be polygenic due to the combined effect of hundreds of points of genetic variance among individuals. One possible variation that affects sports performance is the ACE polymorphism, and another is the ACTN3 polymorphism. The aim of the present study was to analyse the frequency of the polymorphisms of these two genes.

ACE is a key component in the renin-angiotensin-aldosterone system, generating vasoconstrictor angiotensin II and degrading vasodilator kinins (2). ACE is widely expressed in skeletal muscle and other human tissues and may be involved in metabolism during exercise (13-15). A 287-base pair insertion (I) in an Alu intronic region of chromosome 17 has been associated with improved performance in endurance sports, and the corresponding absence of this segment (D) has been associated with improved performance in strength- and power-oriented sports (16). A previous study reported a close similarity in the distribution of genotypes between athletes and control subjects, identifying the following frequencies of genotypes: DD, 30.0 and 29.0%; ID, 47.5 and 49.0%; and II, 22.5 and 22.0%, for athletes and controls respectively (17). In the present study, the DD genotype (which is advantageous for sprinting) was detected at a higher level in the elite athletes (51.35%) than that in the control group (37.84%). The frequency of the II genotype (which has been shown to benefit endurance) was higher in the control group (32.43%) than that in the elite athletes (8.11%). The ID genotype frequency was higher in the elite athletes (40.54%) than that in the control group (29.73%).

The ACTN2 and ACTN3 genes encode the skeletal muscle proteins α-actinin-2 and α-actinin-3, respectively (11).
The first association study to investigate the influence of ACTN3 genotypes on athletic performance was published by Yang et al (11) in 2003. A previous study reported that the frequency of the RR genotype was significantly higher in soccer players (48.3%) than that in controls (28.5%) or endurance athletes (26.5%), and the frequency of the RX genotype was significantly lower in the soccer players (36.7%) compared with that of the controls (53.7%) and endurance athletes (52.0%) (P=0.041). Although there are a notable number of exceptions, elite soccer players tend to have the sprint/power ACTN3 genotype (18). In the present study, the RR (associated with the sprinter property) and XX (associated with the endurance property) genotypes were detected at higher frequencies in elite athletes (10.81 and 35.14%, respectively) than in the controls (2.70 and 10.81, respectively). The frequency of the RX genotype was higher in the control group (86.48%) than that in the elite athletes (54.05%).

The ACE I and ACTN3 X alleles influence speed and power in athletes. Athletes with the ACE I/I and I/D genotypes, as well as the ACTN3 X/X and R/X genotypes, have the potential to achieve improved results in sports requiring power; therefore, analysing the polymorphisms of these genes may be useful in sport type selection (19). However, Myerson et al (4) found that the ACE genotype was not a key determinant of success in the best sprinters worldwide and suggested that the D allele is more common among sprinters. The present study demonstrated that the ACE DD/ACTN3 RR genotype combination is detected at a higher frequency in elite athletes (8.11%) than that in controls (0%). The ACE DD/ACTN3 XX genotype combination was also identified at a higher frequency in the elite group (18.92%) than that in the control group (2.70%). These results are in accordance with those of Eynon et al (20). The frequency of the ACE I allele (present in the ACE II and ACE ID genotypes) and ACTN3 XX genotype combination was higher in the elite athlete group (16.21%) than that in the control group (8.11%). This genotype was therefore identified as enhancing endurance activity performance.

In conclusion, the findings of the present study suggest a novel mechanism involving the renin-angiotensin-aldosterone system, in the response of skeletal muscle fast or slow contraction. However, the aforementioned genotype combination is not associated with the level of performance. Therefore it is difficult to interpret the meaningful results because of the various populations and ethnic groups.

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References