Prevalence of the CYP2D6*10 (C100T), *4 (G1846A), and *14 (G1758A) alleles among Iranians of different ethnicities

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Abstract: The presence of polymorphisms in the CYP2D6 gene may modulate enzyme level and activity, thereby affecting individual responses to pharmacological treatment. Here, we compared the prevalence of the CYP2D6*10, *4, and *14 alleles in an Iranian population of different ethnicities with those of other populations. Allele and genotype frequency distributions of CYP2D6*10 variants and predicted phenotypes including extensive metabolizers, intermediate metabolizers, and poor metabolizers were analysed in blood samples of 300 unrelated healthy individuals in an Iranian population using polymerase chain reaction (PCR)-restriction fragment length polymorphism, PCR-single-strand conformation polymorphism, and direct genomic DNA sequencing. The CYP2D6*4 (G1846A) and *14 (G1758A) allelic frequencies were not detected in different ethnicities, demonstrating the absence of a significant contribution of these alleles in Iranian populations. However, the T/T, C/T, and C/C genotype frequencies of the CYP2D6*10 allele were significantly different (P<0.01) in all Iranian ethnic groups. Additionally, the frequency of the homozygous T/T variant of the CYP2D6*10 allele was significantly high in the Lure (P<0.017) and low in the Kurd (P<0.002) ethnicities. The frequency of the T/T variant of the CYP2D6*10 allele in central Iran was the highest (P<0.001), while the south of Iran had the lowest frequency (P<0.001). The frequency of the C/T variant of the CYP2D6*10 allele was significantly a bit high (P<0.001) in females compared to males, while the frequencies of the T/T variant in females is similar to males, which are 24.4% and 24.3%, respectively. In contrast to absence of the CYP2D6*4 (G1846A) and *14 (G1758A) alleles in Iranian populations of different ethnicities, the prediction of the CYP2D6*10 allele is required in drug research and routine treatment, where the information would be helpful for clinicians to optimize therapy or identify persons at risk of adverse drug reactions before clinical trials. Approximately 39.3% of subjects (24.3% homozygous T/T CYP2D6*10 as poor metabolizers and 15% heterozygous C/T CYP2D6*10 as intermediate metabolizers) had this allele; therefore, the harmful effects of drugs are relatively common among Iranians.

Keywords: pharmacogenetics, polymorphism, cytochrome P450 genes, CYP, Iranian population, antipsychotics, antidepressants

Introduction
Pharmacogenetics is the investigation of different responses of people, based on changes in their genetic structures, towards treatment and drugs. Liver and small intestine microsomal enzymes metabolize the majority of drugs, in which the activity of drugs are determined based on their interaction with enzymatic activity of the cytochrome P450 (CYP) system.1

More than 50 human CYP isozymes have been identified to date,2 and more than 20 are functionally encoded by polymorphic CYP2A6, CYP2C9, CYP2C19,
and CYP2D6 genes.\textsuperscript{2,3} CYP2D6 is of great importance for the metabolism of clinically used drugs, metabolizing approximately 20\%–25\% of these drugs, including antipsychotics, antidepressants, antiarrhythmic and beta-adrenoceptor antagonists (beta blockers).\textsuperscript{4}

The CYP2D6 is a highly polymorphic gene locus with more than 75 allelic variants, and subjects can be classified into poor metabolizers (PM), extensive metabolizers (EM), or ultra-rapid metabolizers of a given CYP2D6 substrate.\textsuperscript{5,6} The CYP2D6*9, CYP2D6*10, CYP2D6*17, and CYP2D6*41 are the most important intermediate metabolizer (IM) alleles;\textsuperscript{7} in which CYP2D6*10 (49.5\%),\textsuperscript{8} CYP2D6*1 (40.1\%–43.5\%),\textsuperscript{9,10} CYP2D6*2 (11.9\%–26.2\%),\textsuperscript{11} and CYP2D6*4 (0.2\%–2.8\%).\textsuperscript{8,10} are the most common CYP2D6 alleles in Asians. Approximately 13\%–50\% of Asian populations may be homozygous for the CYP2D6*10 allele.\textsuperscript{12}

A cytosine-to-thymine substitution (C>T) at nucleotide 100 in the CYP2D6*10 allele, resulting in a modification of proline-to-serine at codon 34, is common among Asian populations\textsuperscript{13} and is associated with reduced metabolism of various CYP2D6 substrates.\textsuperscript{12} There is a highly conserved proline-rich area among microsomal P450s. The proline at codon 34 probably acts as a center between the lipophilic membrane anchor and the active site (heme moiety) of the enzyme. The change of an amino acid in this region may well decrease the activity of the enzyme.\textsuperscript{14,15} The CYP2D6*4 (G1846A) allele results in a splicing defect with nonfunctioning enzyme activity due to replacement of guanine (G) by adenine (A) at position 1,846. This single base substitution in the intron 3/exon 4 boundary causes a shift of the consensus acceptor splice site, yielding a spliced messenger RNA with additional base, thereby resulting in a premature termination codon.\textsuperscript{16} Some studies indicated that the CYP2D6*14 allele may play an important role in contributing to the PM phenotype, in which the CYP2D6*14 allele frequency in Chinese and Japanese populations is approximately 2\%.\textsuperscript{17,18} In this study, the prevalence of the CYP2D6*10, *4, and *14 alleles was investigated in Iranian populations of different ethnicities and in comparison with other countries.

**Materials and methods**

**Sample collection and ethical statement**

Three hundred blood samples were obtained from unrelated and healthy donors from Iranian populations of different ethnicities through the Special Medical Center, Tehran, Iran, including those from 118 Fars, 79 Turk, 42 Mazani (Mazandarani), 37 Lure, and 24 Kurd. Blood samples (2 mL) were taken and collected into tubes with ethylenediaminetetraacetic acid. All the individuals who participated in this study belonged to various ethnicities (Fars, Turk, Mazani, Lure, and Kurd) from different provinces of Iran and were referred for genetic counseling at the Special Medical Center of Tehran between 2006 and 2008. Written informed consent, including consent to participate in the study for genetic analysis and consent to publish, was obtained from the individuals, and the Medical Ethics Committee of the Special Medical Center specifically approved this study. The exclusion criterion for the control healthy group was any history of cancer, metabolic diseases, and nuclear and mitochondrial DNA-related diseases that may affect the DNA.

**DNA extraction and primer sequences**

Genomic DNA was extracted using a MBST kit (salting-out method) from 2 mL of whole blood. Primer sequences for the CYP2D6*10 alleles were ordered as described in a previous study.\textsuperscript{6} The oligonucleotide forward (F) and reverse (R) primers that were used to amplify the CYP2D6*10, *4, and *14 alleles are as follows: F: 5’–TCAACACAGCAGTTCA–3’, R: 5’–CTGTCCTACCCACC–3’ [6]; F: 5’–TGCGCCCTTCGCCAACCCT–3’, R: 5’–TCGCCCTTCAGAGACTCTC–3’\textsuperscript{19} and F: 5’–GTGGATGGTGTTGCTTGAATGCCTT–3’, R: 5’–CAGAGACTCTCGGTCTCCTGCT–3’,\textsuperscript{20} respectively.

**The CYP2D6 metabolism phenotypes**

The classification of CYP2D6 phenotypes is based on the assumption of dominance, in which the predicted phenotype is defined as EM, with one or two active alleles; IM, with two reduced activity alleles or one reduced activity and one inactive allele; and PM, with two inactive alleles.\textsuperscript{16,21}

**Restriction fragment length polymorphism**

Genotyping of the CYP2D6*10 alleles was performed by polymerase chain reaction-restriction fragment length polymorphism (PCR-RFLP). The PCR amplification was carried out with 20–60 ng of genomic DNA, 0.3 U of Taq DNA polymerase, 20 mM of each primers, 10× PCR buffer, 2.5 mM magnesium chloride, and 10 mM deoxynucleotide. The reaction mixture was initially denatured at 95°C for 3 minutes, followed by 35 cycles of 95°C for 1 minute, 52.8°C for 1 minute, 72°C for 2 minutes, and a final extension at 72°C for 7 minutes. The 433 bp amplified fragment was analyzed on 1.5% agarose gel stained with ethidium bromide before restriction digestion. One unit of HphI restriction enzyme was added to each PCR product (5 μL).
and incubated at 37°C for 16 hours. Digested products were analyzed on polyacrylamide gel. The PCR product with a length of 433 bp was subsequently digested with the restriction endonuclease HphI. Only the amplicon of the allele carrying the 100 C>T single-nucleotide polymorphism was cut by HphI between positions 91 and 92 of the CYP2D6 gene, which is between position 258 and 259 of the PCR product (Figure 1). As HphI also cuts at position 362 of the product, irrespective of the amplified allele, the digestion leads to a 362 bp fragment for the wild-type allele and two fragments of 262 bp and 100 bp for the allele carrying the 100 C>T substitution. Additionally, the BSTN1 and MSP1 restriction enzymes were separately added to PCR products in order to cut the amplicon of the CYP2D6*4 and *14 alleles carrying the G1846A and G1758A single-nucleotide polymorphism, respectively.

**Single-strand conformation polymorphism**

The double-stranded, PCR-amplified DNA fragment was denatured before loading on agarose gel. For this purpose, 10 μL of PCR mixture was mixed with 7 μL of freshly prepared denaturation buffer (990 μL of 100% formamide, 10 μL of 1M sodium hydroxide, and a few granules of bromophenol blue) in a 500 μL PCR eppendorf tube. Samples were placed at least 30 minutes in 45°C, and after 23 minutes, 12 μL of the single-strand conformation polymorphism (SSCP) color was added to the sample. After 30 minutes, the samples were analyzed on a 6% acrylamide gel for 16 hours at 110 V. To identify samples, the SSCP gel was stained, and the samples with the C/C, T/T, and C/T genotypes were observed as two, three, and four bands, respectively (Figure 2).

**Sequencing analysis**

The PCR products were examined for specificity using 1.5% agarose gel electrophoresis. Double-stranded DNA automated sequencing was performed using an Applied Biosystems ABI 3100 sequencing machine (Thermo Fisher Scientific, Waltham, MA, USA). All fragments were sequenced in both the F and R directions for confirmation of any nucleotide variation. Sequence variants were analyzed using a FinchTV program (PerkinElmer Inc., Waltham, MA, USA) (Figure 3).

**Statistical analysis**

The statistical analysis was conducted using SPSS 17.0 (for Windows) to perform the chi-square and 95% confidence interval (CI) tests to calculate the CYP2D6*10 frequencies in different ethnicities and different geographical regions of Iran. A P-value of <0.01 was considered statistically significant.

**Results**

Allele and genotype frequency distribution of the CYP2D6*10, *4 (G1846A), and *14 (G1758A) variants was analyzed in blood samples of 300 unrelated healthy individuals in an Iranian population of different ethnicities. The allele frequencies of the homozygous (T/T), heterozygous (C/T), and wild-type (C/C) variants of CYP2D6*10 were significantly different (χ²=90.02, P<0.01) in all Iranian ethnic

**Notes:** The two, three, and four bands in each lane represent the homozygous wild-type (C/C), homozygous (T/T), and heterozygous (C/T) alleles of the CYP2D6*10, respectively; Lanes 1 and 2 represent the homozygous C/T genotype (four fragments), lane 3 represents the homozygous T/T genotype (three fragments), while lane 4 represents the wild-type C/C genotype (two fragments). 

**Abbreviation:** PCR-RFLP, polymerase chain reaction-restriction fragment length polymorphism.

**Figure 1** PCR-RFLP analysis of the CYP2D6*10 allele using the Hphl enzyme.

**Notes:** Lane 1 indicates the PCR product; This enzyme generated four fragments of 362 bp, 262 bp, 100 bp, and 71 bp (lane 2) in the heterozygous C/T genotype; three fragments of 262 bp, 100 bp, and 71 bp (lane 3) in the homozgyous T/T genotype; and two fragments of 362 bp and 71 bp (lane 4) in length in the wild-type C/C genotype.

**Abbreviation:** PCR-RFLP, polymerase chain reaction-restriction fragment length polymorphism.

**Figure 2** PCR-SSCP analysis of the CYP2D6*10.

**Notes:** The two, three, and four bands in each lane represent the homozygous wild-type (C/C), homozygous (T/T), and heterozygous (C/T) alleles of the CYP2D6*10, respectively; Lanes 1 and 2 represent the heterozygous C/T genotype (four fragments), lane 3 represents the homozygous T/T genotype (three fragments), while lane 4 represents the wild-type C/C genotype (two fragments). 

**Abbreviation:** PCR-SSCP, polymerase chain reaction-single-strand conformation polymorphism.
groups. Genotype combinations were used to categorize CYP2D6 metabolism phenotypes as PM, IM, and EM (Table 1). Additionally, the frequency of the homozygous (T/T) variant of *CYP2D6* was significantly high in the Lure ($\chi^2=8.162$, $P<0.017$) and low in the Kurd ($\chi^2=12.25$, $P<0.002$) ethnicities. However, the frequency of the homozygous (T/T) variant of *CYP2D6* in ethnic Fars was similar to that of ethnic Mazani (Table 1). The frequency of the heterozygous (C/T) variant of *CYP2D6* was significantly high in Mazani ($\chi^2=10.857$, $P<0.004$) and low in Kurds ($\chi^2=12.25$, $P<0.002$). Moreover, the frequency of the wild-type (C/C) variant of *CYP2D6* was highest in Kurds, yet lowest in Lures (Table 1). The results indicate that the frequency of the homozygous (T/T) variant of *CYP2D6* in central Iran was significantly the highest ($\chi^2=15.354$, $P<0.001$), while the south of Iran had the lowest frequency ($\chi^2=23.164$, $P<0.001$). The frequency of the heterozygous (C/T) variant of *CYP2D6* allele in the east of Iran had the highest frequency, while the west had the lowest frequency (Figure 4; Table 1). The frequency of the C/T variant of the *CYP2D6* allele was significantly a bit higher ($P<0.001$) in females compared to males, while the frequencies of the T/T variant in females was similar to males, which are 24.4% and 24.3%, respectively (Table 1). However, the

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**Figure 3** PCR-sequencing of the *CYP2D6* allele (C100T).

Notes: The chromatogram shows the mutant allele (A) compared to the wild-type allele as highlighted in blue color (B). The arrows mark the polymorphism.

Abbreviation: PCR, polymerase chain reaction.
CYP2D6*4 (G1846A) and *14 (G1758A) allelic frequencies were not detected in different ethnicities, demonstrating the absence of a significant contribution of these alleles in Iranian populations. Additionally, the CYP2D6*10 allele was directly detected using the PCR-RFLP and -SSCP. The PCR-RFLP analysis of the CYP2D6*10 allele using the HphI enzyme indicates the heterozygous C/T, homozygous T/T, and wild-type C/C genotypes (Figure 1). The PCR-SSCP analysis demonstrates the heterozygous C/T, homozygous T/T, and wild-type C/C genotypes with four, three, and two fragments, respectively (Figure 2).

**Discussion**

CYP2D6 is a clinically important enzyme that metabolizes numerous therapeutic drugs. Among the alleles of CYP2D6, CYP2D6*10 (C100T) is the most common allele in the Asian population, with a frequency of between 30% and 69.2%, where 13%–49% of the aforementioned population are homozygous for this allele. Accordingly, the majority of the Asian population is at risk of either harmful effects or adverse reactions towards medication. Thus, considering the effects of the CYP2D6*10 allele on drug metabolism in Iran, along with allele genotyping, including that of CYP2D6*wt*/wt, CYP2D6*wt*/10, and CYP2D6*10/*10, may prevent the harmful effects of drugs and drug interactions, thereby reducing mortality risk in Iranian populations.

The CYP2D6*10 “C100T” mutation in patients leads to reduced enzyme activity and decrease of appropriate response to drug treatments, such as tamoxifen, codeine, antidepressants, and antipsychotics. Additionally, there has been some research linking the CYP2D6 alleles and drug metabolism in the Iranian population, but study of the CYP2D6*10 allele regarding its significant role in drug metabolism in Asian populations have been neglected. Therefore, genotyping of the CYP2D6*10 allele, including CYP2D6*wt*/wt, CYP2D6*wt*/10, and CYP2D6*10/*10, in different ethnicities including Lure, Turk, Fars, Kurd, and Mazani is considered as an important determinant for the prevention of the harmful effects of drugs and drug interactions, resulting in better survival and reduced mortality risk.

The CYP2D6 allele frequency in our Iranian population with different ethnicities including Lure, Turk, Fars, Kurd, and Mazani were compared with the frequencies seen in other ethnic populations. The study by Kouhi et al. showed that the prevalence of CYP2D6*10 was significant only in the eastern part (Tabriz) of Azerbaijan (Turk). However, in our study, the samples were selected from both the western and eastern parts of Azerbaijan (Turk), indicating no significant difference in Turk ethnicity. The CYP2D6*10 allele was found to have a frequency of 0.319 in the Iranian population (Figure 5), which is higher than that of South Indian and Malaysian populations, but not as high as what was reported in other populations such as the Japanese, and Chinese.

There are four variations including C100T, G1758A, C2850T, and G4180C in the CYP2D6*14 allele, in which identification of the G1758A polymorphism is exclusively used for the CYP2D6*14 genotype determination. In our
Figure 4 Geographical distribution of the CYP2D6*10 genotypes among 300 normal Iranian individuals of different ethnicities.

Notes: The T/T, C/T, and C/C genotype frequencies of CYP2D6*10 were significantly different ($\chi^2$=90.02, P<0.01): 24.3%, 15%, and 60.7%, respectively; The results indicate that the frequency of the homozygous T/T variant of CYP2D6*10 in the central Iran was the highest ($\chi^2$=15.354, P<0.001), while the south of Iran had the lowest frequency ($\chi^2$=23.164, P<0.001); The frequency of the heterozygous C/T variant of CYP2D6*10 allele is highest in the east and lowest in the west; The green, pink, yellow, blue, and red colors indicate the north, south, east, west, and center of Iran, respectively.

Figure 5 Frequency of the CYP2D6*10 allele in different populations.
study, the CYP2D6*4 (G1846A) and *14 (G1758A) allelic frequencies were not identified in different ethnicities including Fars, Turk, Lure, Kurd, and Mazani, demonstrating the absence of a significant contribution of these alleles in Iranian populations compare to other populations (Figure 6).

In conclusion, this study demonstrates the absence of the CYP2D6*4 (G1846A) and *14 (G1758A) alleles, whereas approximately 24% of the Iranian population has the homoygous T/T CYP2D6*10, suggesting susceptibility to adverse or poor drug responses, which may lead to increase in mortality risk. It is clinically important to identify those individuals who probably have altered pharmacokinetics for CYP2D6 substrates in order to avoid adverse drug reactions. CYP2D6 polymorphism analysis in different ethnicities indicates a very useful forensic tool toward personalized medicine. These data also allow comparison of pharmacogenetic variation between populations with different ethnicities. Therefore, the determination of the presence of this allele in individuals via genetic tests is a basic step to prevent the harmful effects of drugs.

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Disclosure

The authors report no conflicts of interest in this work.

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