

Full length article

Distributive characteristics of Ser49Gly and Gly389Arg genetic polymorphisms of β_1 -adrenoceptor in Chinese Han and Dai populations¹

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Key words

β_1 -adrenoceptor; genetic polymorphism; Ser49Gly; Gly389Arg

¹ Project supported by the National Natural Science Foundation of China (No 30070880), and the Sponsorship for Outstanding Teachers Program, Ministry of Education, China (No 2002-350).

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Received 2005-07-24

Accepted 2005-10-10

doi: 10.1111/j.1745-7254.2006.00248.x

Abstract

Aim: Genetic polymorphisms causing Ser49Gly and Gly389Arg mutants of β_1 -adrenoceptor may result in significant changes in the function of this receptor. The aim of the present study was to investigate the frequencies of the Ser49Gly and Gly389Arg mutant alleles in healthy Chinese populations and to investigate the differences between 2 Chinese ethnic groups (Han and Dai populations) with respect to the frequencies of these alleles. **Methods:** A total of 225 Han Chinese and 175 Dai Chinese unrelated healthy volunteers were recruited for this study. Genomic DNA was extracted from peripheral blood leukocytes by using a standard manual chloroform-phenol extraction. Fragments spanning the 2 polymorphisms were amplified by using polymerase chain reaction with template genomic DNA and relevant primers. The DNA products including the polymorphic loci were subjected to restriction endonuclease digestion with *Eco*0109I and *Bcg*I. Digested fragments were detected with an ultraviolet detector after electrophoresis (100 V for approximately 1.5 h). **Results:** The frequencies of the Gly49 and Arg389 alleles were, respectively, 16.2% and 76.4% in the Han population and 14.6% and 75.7% in the Dai population. **Conclusion:** The polymorphisms causing the Ser49Gly and Gly389Arg mutations of the β_1 -adrenoceptor existed in both healthy Han and Dai Chinese populations. The frequencies of the Ser49Gly and Gly389Arg mutant alleles were not significantly different in the Han and Dai populations. However, the frequency of the Gly389 variant seems to be significantly lower in these 2 populations than in an African-American population.

Introduction

Human β -adrenergic receptor is a member of the 7-transmembrane superfamily of G protein-coupled receptors. β_1 -adrenoceptor is encoded by an intronless gene located on chromosome 10q24–26, and it was firstly cloned and sequenced in 1987^[1–3]. In human heart, β_1 -adrenoceptor is the predominant receptor that modulates cardiac inotropy and chronotropy in response to endogenous catecholamines through coupling to G_s to activate adenylyl cyclase^[4,5]. Recently, Podlowski *et al* identified 7 different single-nucleotide polymorphisms (SNP) in the β_1 -adrenoceptor gene by using polymerase chain reaction-single strand conformation

polymorphism (PCR-SSCP) analysis, and found that each SNP led to an amino acid exchange^[6]. Two of these SNP have also been reported by previous studies, and are thought to be of functional significance. One of the 2 functionally important SNP is located at the amino-terminus: an A→G exchange at 145 bp (A145G) results in an amino acid substitution of Ser by Gly at residue 49 (Ser49Gly). A previous study confirmed that the Ser to Gly substitution affected agonist-promoted trafficking, and reported that the Gly49 polymorphic receptor had enhanced agonist-promoted downregulation^[7]. It had also been reported that the Gly49 allele frequency was approximately 0.13–0.15 in a Caucasian American population; however, racial differences in the dis-

tribution of the Gly49 allele have not yet been reported. A previous study showed that a genetic polymorphism in the β_1 -adrenoceptor gene existed in patients with idiopathic dilated cardiomyopathy (IDCM) and congestive heart failure^[8].

A G→C exchange at 1165 bp in the β_1 -adrenoceptor gene causes amino acid substitution of Gly by Arg at residue 389 (Gly389Arg). The Gly389Arg polymorphism occurs within the intracellular cytoplasmic tail near the 7-transmembrane-spanning segment of the β_1 -adrenoceptor. A previous *in vitro* study confirmed that the Gly to Arg substitution markedly altered the G-protein coupling of the β_1 -adrenoceptor molecule, and that the level of cAMP generated by β_1 -adrenoceptor activation following exposure to an agonist differed by about 3-fold for the Arg389 and Gly389 variants^[9]. It seems that there are racial differences in the frequency of alleles at codon 389, the frequencies of the Arg389 allele in Caucasian Americans and African-Americans were 0.73 and 0.58, respectively^[10,11]. This allele may contribute to individual and interethnic differences in drug responses, primarily through β_1 -adrenoceptor modulation.

China is a multicultural country; the population comprises 55 ethnic minorities in addition to the Han majority. Each of these ethnic groups has a unique genetic background. The Dai group is the 17-largest ethnic group in China, comprising more than one million people, who live chiefly in the west of Yunnan province, in South-western China^[12]. Our previous studies showed that the incidence of drug metabolizing enzyme polymorphisms varies in different ethnic groups^[13]. It was therefore interesting to determine whether the incidence of β_1 -adrenoceptor polymorphisms were concordant in 2 different ethnic populations in the same country.

β_1 -Adrenoceptor is an important target of many drugs and endogenous substances. A better understanding of individual and interethnic differences with respect to β_1 -adrenoceptor polymorphisms may help us to explain intraindividual differences in drug response and disease susceptibility. Therefore, in the present study we investigated the frequencies of the main mutants of the β_1 -adrenoceptor gene in Han and Dai Chinese populations. Because we consider that polymerase chain reaction-restriction fragment length polymorphism (PCR-RFLP) analysis is currently one of the most reliable methods for population genotyping, PCR-RFLP methods for Ser49Gly and Gly389Arg genotyping of the β_1 -adrenoceptor gene were used.

Materials and methods

Subjects The study protocol was approved by the Eth-

ics Committee of Xiang-Ya School of Medicine, Central South University. A total of 225 Han Chinese (123 men, and 102 women) living in Hu-nan province and 175 Dai Chinese living in Yunnan province (89 men, and 86 women) were recruited for the present study. The subjects were unrelated healthy volunteers aged 18–21 years, and all participated after giving their written informed consent. All of the Han subjects were students from Xiang-Ya School of Medicine, Central South University, and the Dai subjects were students from the Dali Nationality School, Yunnan province. None of the subjects had any abnormalities according to their medical records, routine laboratory tests, physical examinations, or electrocardiography. Samples (5 mL per subject) of peripheral vein blood were collected for genotyping analysis (Table 1).

Table 1. Clinical characteristics of the Han and Dai Chinese populations.

Characteristic	Ethnic group		P-value
	Han (n=225)	Dai (n=175)	
Sex (M/F)	123/102	89/86	0.262
Age (years)	19.5±0.4	20.2±0.3	0.138
Height (cm)	167.7±2.1	168.4±2.8	0.183
Weight (kg)	61.4±3.1	62.1±2.9	0.906
BMI (kg/m ²)	22.5±1.1	21.1±1.1	0.237
HR (beat/min)	72.6±1.9	68.0±1.6	0.084
SBP (mmHg)	114.1±3.8	111.3±3.4	0.219
DBP (mmHg)	75.8±2.5	72.2±2.7	0.443

BMI, body mass index; HR, heart rate; SBP, systolic blood pressure; DBP, diastolic blood pressure.

Genotyping procedures for β_1 -adrenergic receptor Genomic DNA was extracted from peripheral lymphocytes with phenol-chloroform followed by ethanol precipitation^[14]. Genotyping analysis was carried out by using the PCR-RFLP assay. The PCR procedure for the β_1 -adrenergic receptor gene was performed as described previously with minor modifications^[15]. For the Ser49Gly locus, we used the following primer pairs: the sense primer P1, 5'-CCGGGCTTCTGGGGTGTTC-3', and the antisense primer P2, 5'-GGCGAGGTGATGGCGAGGTAGC-3'. The Gly389Arg polymorphic locus was amplified by using the sense primer P3, 5'-CATCATGGGCGTCTTCACGC-3', and the antisense primer P4, 5'-TGGGCTTCGAGTTCACCTGC-3'. The amplified DNA fragments including the Ser49Gly or Gly389Arg polymorphic sites were separately digested with *Eco*0109I (TaKaRa Biotech, China) or *Bcg*I (England Biolabs, Beverly, USA) at 37 °C for 8 h.

The different patterns produced by the digested fragments were visualized on 2% agarose gels stained with ethidium bromide. Following the amplifications, some of the PCR product was purified and sequenced to confirm the accuracy of the RFLP assay.

Ser49Gly and Gly389Arg point mutations The Ser49Gly point mutation gave rise to an *Eco*0109I cleavage site. The PCR product of the Gly49 allele contained a unique site for restriction by *Eco*0109I. The PCR product of the Gly49 allele contained a unique site for restriction by *Eco*0109I. The Ser49Gly mutant allele had 3 bands, 564 bp, 345 bp, and 219 bp, which became apparent when digested fragments were detected with an ultraviolet detector (Figure 1). The Gly389Arg substitution gave rise to a *Bcg*I cleavage site. The Gly389Arg mutant had 3 bands, 530 bp, 342 bp, and 154 bp. *Bcg*I cleaves twice, excising its recognition site, which accounts for the 34 bp discrepancy in the fragments generated (Figure 2). When the PCR-RFLP genotyping assay was repeated for randomly selected samples, the result of the second genotyping of each sample was identical to the first in every case, thus verifying the reproducibility of the genotyping results.

Statistical analysis Data analysis was performed using SPSS (version 10.0 for Windows, SPSS, Chicago, IL, USA). A two-tailed value of $P < 0.05$ was considered statistically significant. The frequencies of each allele polymorphism were calculated based on the observed number of different alleles (49Ser or 49Gly, or 389Gly or 389Arg) in each ethnic group. Hardy-Weinberg equilibrium and possible differences in allele frequencies between the 2 populations were tested by using the χ^2 -test.

Results

In the present study, we found that the 2 single nucleotide polymorphisms causing the Ser49Gly and Gly389Arg mutants of the β_1 -adrenoceptor existed in both the healthy Han and Dai Chinese populations.

The allele and genotype frequencies for the β_1 -adrenoceptor Ser49Gly and Gly389Arg polymorphisms in Han and Dai populations are summarized in Table 2. The allelic frequencies for Gly49 and Arg389 were 16.2% and 76.4% in the Han population and 14.6% and 75.7% in the Dai population, respectively. The allele and genotype frequencies for the 2 different populations were in Hardy-Weinberg equilibrium as tested by comparisons of the frequencies found versus the frequencies expected on the basis of the proportion represented by the more frequently occurring allele. There was no difference between the Han and Dai populations with respect to either of the allele frequencies. Compared with data in the literature regarding the 2 polymorphisms in healthy

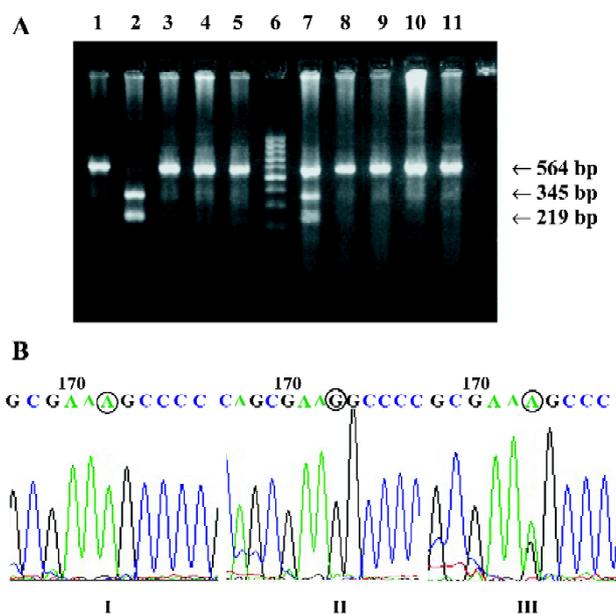


Figure 1. (A) RFLP analysis of the Ser49Gly polymorphism of the β_1 -adrenergic receptor gene. *Eco*0109I-digested fragments of PCR-amplified products of the β_1 -adrenergic receptor gene were separated on a 2% agarose gel and stained with ethidium bromide. Lane 6: 100 bp DNA ladder; lanes 1, 3–5, 8–10: Ser49 homozygotes, which contain only a single 564 bp fragment; lane 2: Gly49 homozygote, which has no 530 bp fragment, but rather two 345 and 219 bp fragments; lane 7: heterozygote, which has all 3 fragments (564, 345, and 219 bp). (B) Sequence of the β_1 -adrenergic receptor Ser49Gly mutant. I, Ser49Ser receptor; II, Gly49Gly receptor, III, Ser49Gly receptor.

subjects from different ethnic groups, significant interethnic differences in the incidence of the Arg389 polymorphism were found between a Chinese and an African American population; however, the frequencies of the Ser49Gly polymorphism were not significantly different in these 2 populations^[11].

Discussion

The β_1 -adrenoceptor is an important target for many drugs and endogenous substances^[16]. In the present study, we carried out the first investigation of the frequencies of the Ser49Gly and Gly389Arg genetic polymorphisms of β_1 -adrenoceptor in Han and Dai Chinese populations and found the frequencies for both polymorphisms were not significantly different in the 2 populations; however, the frequency of the Gly389 variant appeared to be significantly lower in the Han and Dai Chinese populations than in an African-American population.

In the present study, we found that the allelic frequencies of Gly49 and Arg389 were approximately 14.6%–16.2%

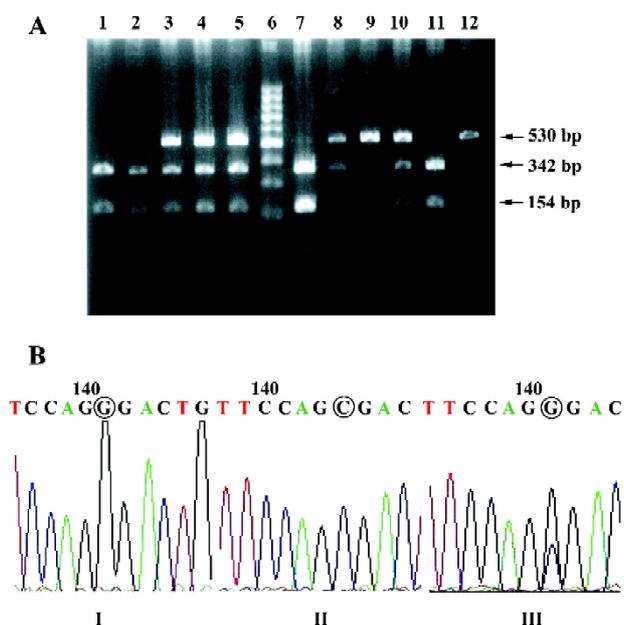


Figure 2. (A) RFLP analysis of the Gly389Arg polymorphism of the β_1 -adrenergic receptor gene. *BcgI*-digested fragments of PCR-amplified products of the β_1 -adrenergic receptor gene were separated on a 2% agarose gel and stained with ethidium bromide. Lanes 1–2, 7, 11: Arg389 homozygotes, which have 342 and 154 bp fragments; lanes 9, 12: Gly389 homozygotes, with only a single 530 bp fragment; lanes 3–5, 8, 10: Gly389Arg heterozygotes, which have all 3 fragments (536, 342, and 154 bp). *BcgI* cleaves twice, excising its recognition site, which accounts for the 34 bp discrepancy in the fragments generated. (B) Sequence of the β_1 -adrenergic receptor Gly389Arg mutant. I, Gly389Gly receptor; II, Arg389Arg receptor; III, Gly389Arg receptor.

and 75.7%–76.4% in the combined Han and Dai populations, respectively. Gly389 is considered to be the wild type, and was first cloned by Frielle *et al* as a Gly389-containing receptor^[1].

The Dai ethnic group is one of many minority groups in China who have unique characteristics with respect to genetic make-up, culture, diet and environment^[12]. Previous studies carried out by our laboratory have found that interethnic differences with respect to polymorphisms in drug metabolizing enzymes such as CYP2C19 exist between Han and Dai populations. He *et al* reported that the frequency of the CYP2C19*3 allele in a Dai population was significantly lower than that in a Han population^[13]. In the present study, we found that there was no significant difference in the frequencies of the β_1 -adrenoceptor Ser49Gly and Gly389Arg polymorphisms between the two ethnic groups. An explanation for the difference between the 2 enzymes with respect to the level of polymorphism in different populations is that

Table 2. Genotype and allele frequencies of β_1 -adrenoceptor Ser49Gly and Gly389Arg polymorphisms in Han and Dai Chinese populations.

Polymorphism	Ethnic group		χ^2	P
	Han (n=225)	Dai (n=175)		
Ser49Gly genotype				
49Ser/Ser	156 (69.3%)	126 (72.0%)	0.509	0.775
49Ser/Gly	65 (28.9%)	47 (26.9%)		
49Gly/Gly	4 (1.8%)	2 (1.1%)		
Ser49Gly allele frequency				
49Ser	377 (83.8%)	299 (85.4%)	0.410	0.522
49Gly	73 (16.2%)	51 (14.6%)		
Gly389Arg genotype				
389Gly/Gly	10 (4.4%)	9 (5.1%)	0.111	0.946
389Gly/Arg	86 (38.2%)	67 (38.3%)		
389Arg/Arg	129 (57.3%)	99 (56.6%)		
Gly389Arg allele frequency				
389Gly	106 (23.6%)	85 (24.3%)	0.058	0.810
389Arg	344 (76.4%)	265 (75.7%)		

Genotype and allele distribution are indicated as absolute values, with percentages in parentheses.

β_1 -adrenoceptor may well be more conserved throughout different populations than drug metabolizing enzymes. Many drugs that have effects mediated by β_1 -adrenoceptor have obvious interethnic differences in their effects^[17]. If such a difference was also found between Han and Dai populations, the differences would be unlikely to be due to β_1 -adrenoceptor polymorphisms, but would rather be due to genetic variations in drug-metabolizing enzymes and the effects of environmental factors.

A previous study reported that the incidence of the Ser49Gly amino acid substitution was 16.2% in patients with IDCIM, but zero in a healthy population^[6]. In contrast, we found that the frequency of the Gly49 allele was approximately 14.6%–16.2% in healthy subjects, which is consistent with the results of other studies, in which the frequency of the Gly49 allele was found to be 2%–15% in healthy subjects^[8,15,18]. Because one study in which the Gly49 allele was only detected in patients with IDCIM used only a small sample size (37 patients), the resulting in low statistical power might have produced a false result. However, several other studies that used a large sample size have also found that there was a higher frequency of the Gly49 allele in patients with IDCIM relative to healthy subjects^[6,8,15].

An *in vitro* mutagenesis study of the codon 389 poly-

morphism revealed that the Arg389 receptor form had nearly two-fold greater basal and 3-fold greater agonist-mediated adenylyl cyclase activity^[9], which suggests that the Gly389 mutation produces a hypoactive β_1 -adrenoceptor, and that a higher frequency of the Gly389 allele could contribute to decreased sensitivity to β -blockers in African-Americans. However, we found no ethnic differences in the frequency of the β_1 -adrenoceptor Gly389Arg polymorphism between Han and Dai Chinese populations in the present study, indicating that this polymorphism is unlikely to account for any ethnic differences in sensitivity to β -blockers between these two populations.

In conclusion, we found that the β_1 -adrenoceptor Ser49Gly and Gly389Arg mutations both existed in both Han and Dai Chinese populations, and that the frequencies of the two polymorphisms were not significantly different in these two ethnic groups.

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