The association between an endothelial nitric oxide synthase gene polymorphism and coronary heart disease in young people and the underlying mechanism

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Abstract. With the development of molecular biological technology, the association between genes and diseases has drawn increasing attention of researchers; the endothelial nitric oxide synthase (eNOS) gene has been reported to be a candidate gene for cardiovascular disease (CHD). The present study aimed to investigate the association between a polymorphism of eNOS and the risk of CHD in young people (≤40 years old), in addition to the underlying mechanism. A total of 234 cases of CHD in young individuals were collected as the CHD group and 228 cases of healthy individuals as the control group. Peripheral blood was collected and the genotype of the eNOS G894T polymorphism was identified by polymerase chain reaction-restriction fragment length polymorphism, the gene frequency was calculated and the distributions of genotype and allele frequency between the two groups were compared. Bioinformatics tools were employed to analyze the differences in the local protein structures of the eNOS G894T polymorphism and the biological mechanism was preliminary discussed. The results demonstrated that there were significant differences in the distribution of genotype frequency and allele frequency between the two groups. Bioinformatics tools were employed to analyze the differences in the local protein structures of the eNOS G894T polymorphism and the biological mechanism was preliminary discussed. The results demonstrated that there were significant differences in the distribution of genotype frequency and allele frequency of the eNOS G894T gene polymorphism between the CHD group and control group (P<0.05). The risk of CHD in GT and TT genotypes were higher compared with the GG genotype (P<0.05). The G894T polymorphism led to Glu298Asp mutation of encoded protein, which is within the active site of eNOS, and partial structures of the protein were converted from random coil to α-helix. In conclusion, the eNOS G894T gene polymorphism was associated with the occurrence and development of CHD in young people. The potential mechanism is that the G894T polymorphism leads to altered protein structure, which affects the function of eNOS in generating nitric oxide and cardiovascular diastole. The results of the present study suggested a potential target gene for the prevention and treatment of CHD in young people (≤40 years old).

Introduction

Coronary heart disease (CHD) is a type of cardiovascular disease causing by coronary artery atherosclerosis and plaque rupture, leading to the arterial stenosis or occlusion, then resulting in myocardial ischemia or necrosis, which is among the top ten causes of human mortality. At present, the treatment methods of CHD include: Surgical treatment, interventional therapy (stent implantation) and medication (nitric acid ester, β-blocker, antithrombotic and statins); however, medication is the basis of all treatment (1). With the improvement of the living standards, the incidence and mortality rates of CHD has increased rapidly, particularly within young people (≤40 years old) in China (2,3). The earlier incidence of CHD is not only threat to human health, but may burden a patient's families and society; CHD has gained increasing attention of researchers (4). It is well-known that smoking, drinking, obesity, hypertension and metabolic syndrome are risk factors for CHD. For smokers, nicotine in tobacco may damage the vascular endothelium, resulting in the vasospasm and plaques rupture, accelerating the onset of coronary atherosclerosis (5). In addition, previous studies have demonstrated that the functional disturbance of endothelial cells and early-stage atherosclerosis may be caused by reduced levels of nitric oxide (NO), nitric oxide has an important role in the pathogenesis of CHD (6-8). Endothelial NO synthase (eNOS) is the enzyme responsible for the generation of NO in endothelial cell (9-11). Various studies have investigated the association between the eNOS G894T polymorphism and the risks of CHD (12-15). However, to the best of our knowledge, the association
between eNOS and the risk of CHD in young people is yet to be established. Therefore, in the present study, the association between the eNOS G894T polymorphism and CHD in young individuals, and the influence of the polymorphism on the protein structure and function, were analyzed and the potential biological mechanisms were preliminarily investigated by a series of bioinformatics analyses.

**Materials and methods**

**Study subjects.** A total of 234 cases of young patients with CHD, who received treatment between August 2013 and May 2016 at the Department of Cardiology, The First Affiliated Hospital of Zhengzhou University (Zhengzhou, China) were included in the CHD group, which comprised 198 males and 36 females between the ages of 18 and 40 years old with a mean age of 34.2±3.7 years old. All cases were diagnosed with CHD through a coronary angiogram and complied with the following CHD diagnostic criteria specified by the World Health Organization (16): Exhibited symptoms typical of angina; an electrocardiogram identified the previous occurrence of a myocardial infarction; and confirmation by angiography, which is considered the ‘gold standard’ for the assessment of coronary artery disease. A comprehensive individualized treatment program should be applied to suit the circumstances of each patient. A total of 228 healthy individuals between the ages of 18 and 40 years old who visited the hospital for physical examination and voluntarily agreed to participate in the study during the same period were randomly collected as the control group, which comprised 195 males and 33 females with a mean age of 35.7±2.6 years old. All of the participants lived in the Henan province of China and had no blood relationship with each other. The present study was reviewed and approved by the Ethical Inspection Committee of Zhengzhou University and participants in the CHD and control groups volunteered and signed the informed consent.

**Identifying genotypes by polymerase chain reaction-restriction fragment length polymorphism (PCR-RFLP).** Peripheral blood (4 ml) was collected from each participant, 0.6 mg EDTA-Na$_2$ was used as an anticoagulant (final concentration was 1.5 g/l) and the genomic DNA was extracted using a Genelute™ blood genomic DNA kit (Sigma-Aldrich; Merck KGaA, Darmstadt, Germany) according to the manufacturer’s protocols. The diagnostic primer for the genotype of the G894T polymorphism was synthesized by Sangon Biotech Company (Shanghai, China): Sense, 5'-GAGATGAAGGCAGAGCACTG-3'; and anti-sense, 5'-TCCATCCCCACCTGCAAT-3'. The PCR reaction system (25 µl) constituted: Genomic DNA 1.0 µl, 10x buffer 2.5 µl, dNTP Mix 2.0 µl, sense primer 0.2 µl, anti-sense primer 0.2 µl, Taq DNA polymerase 0.1 µl and deionized water 19 µl. The amplification procedure: pre-denaturation at 95°C for 2 min; 35 cycles of denaturation at 95°C for 30 sec, annealing at 58°C for 30 sec and extension for 30 sec at 72°C and final extension at 72°C for 5 min. The PCR products were purified with Gel PCR purification kit (Thermo Fisher Scientific, Inc., Waltham, MA, USA) according to the manufacturer’s protocols. PCR-amplified specific fragments of eNOS were obtained. Amplified products (10 µl) were digested with 0.5 µl restriction endonuclease Eco24I (Ban II) (Takara, Japan) at 37°C for 10 h. The results of enzyme digestion were performed by agarose gel electrophoresis (Fig. 1) and the results demonstrated that wild-type homozygote GG contained 169 and 94 bp fragments, variant homozygote TT contained a single 263 bp fragment and heterozygote GT contained 263 and 169 and 94 bp fragments. The G849T polymorphism from individuals of the present study had been sequence by PCR-RFLP; in the GG genotype, the G849T locus was at a guanine nucleotide, which could be digested by Eco24I. Within the TT genotype, the G849T locus was at a thymine nucleotide; however, within the GT type, the G849T locus was a guanine or thymine nucleotide.

**Bioinformatics analysis on the polymorphism site.** Alterations in the amino acid sequences of the eNOS G894T gene polymorphism were analyzed by the Single Nucleotide Polymorphisms and open reading frame databases (National Center for Biotechnology Information (NCBI); https://www.ncbi.nlm.nih.gov/; https://www.ncbi.nlm.nih.gov/orffinder/, respectively), and alterations in the protein structure prior to and following the amino acid sequence mutation were analyzed using the protein modeling tool, QUARK (17). Amino acid sequences consisting of 20 amino acids around the G894T polymorphism site were entered into the QUARK tool to perform local structural modeling of the protein, and structural characteristics prior to and following the mutation were analyzed and compared by Cn3D 4.3.1 viewer software (NCBI; http://www.ncbi.nlm.nih.gov). Characteristics of the eNOS structural domain were analyzed by SMART v7 (18) and the NCBI Conserved Domain Database (CDD) (19), and the potential biological functions of the region where the polymorphism sites were located was discussed. In addition, the potential eNOS signaling pathway and the effects of structural changes on the pathway activity were analyzed by using the Kyoto Encyclopedia of Genes and Genomes (KEGG) pathway database (20) (http://www.genome.jp/kegg/pathway.html).

**Statistical analysis.** All data were analyzed by SPSS 19.0 (IBM Corp., Armonk, NY, USA). Data are presented as the mean ± standard deviation, and the comparisons between two groups were performed by Student's t-test. The comparison of enumeration data in two groups (including sex, smoking, hypertension and genotype, Tables I, II and III) was performed.
by Chi-squared (χ²) test. P<0.05 was considered to indicate a statistically significant difference. Hardy-Weinberg equilibrium was used to analyze whether samples in each group were representative of the population and P>0.05 was considered to indicate that the sample was representative. The differences in the distribution of genotype and alleles between the two groups were compared by the χ² test and binary logistic regression was performed to analyze the association between genotype and disease risk, and the odds ratio (OR) value was presented to indicate the relative risk.

Results

Subject characteristics. The characteristics of the control and CHD groups are presented in Table I. There were no statistically significant differences in age, sex, body mass index, smoking, diabetes, hypertension or family history of CHD between the CHD and control groups (P>0.05).

Distribution of genotype and allele frequency of the G894T site. As demonstrated in Table II, the distribution of genotype in the control and CHD groups conformed to the Hardy-Weinberg equilibrium (P>0.05), which indicates that the samples were typically representative. The differences in the distribution of genotype and alleles between the two groups were compared by the χ² test and logistic regression was performed to analyze the association between genotype and disease risk, and the odds ratio (OR) value was presented to indicate the relative risk.

Partial structural modeling of the G894T locus. The eNOS G894T gene polymorphism leads to a change in the amino acid located at the 289th amino acid position, changing from glutamic acid (Glu/E) to aspartic acid (Asp/D), and the result of modeling is presented in Fig. 2 as obtained from QUARK (14) and the Cn3D 4.3.1 viewer software (http://www.ncbi.nlm.nih.gov). When the 289th amino acid was changed from Glu to Asp, the tertiary structure of the short-peptide changed from a random coil to an α-helix, indicating that the local structure of the protein in different genotypes was different, which may result in alterations to the protein function.

Analysis of structural domain. The structural domain of eNOS was analyzed by SMART and results are presented in Fig. 3. The G894T site was located in the low complexity region at the 287-321th amino acid of eNOS (Fig. 3A), which indicated that the region may be involved in flexible binding associated with specific functions and important in determining binding properties and biological roles (21). Furthermore, CDD analysis demonstrated that G894T was located in the active region of the eNOS enzyme, which binds to the substrate L-arginine, zinc, the cofactor heme, tetrahydrobiopterin and the C-terminal electron supplying reductase region. Therefore, it may be hypothesized that the G894T polymorphism may group, while the risk of CHD in TT and GT + TT groups was 2.075 and 1.889 times the risk in the GG group, respectively (P<0.05; Table III). Furthermore, the risk of CHD associated with the eNOS 894T allele was 1.749 times the risk associated with the eNOS 894G allele (P<0.05; Table III).

Table I. Comparison of subject characteristics between the CHD and control groups.

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>CHD group (n=234)</th>
<th>Control group (n=228)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, years</td>
<td>34.2±3.7</td>
<td>35.7±2.6</td>
<td>0.73</td>
</tr>
<tr>
<td>Sex</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>198</td>
<td>195</td>
<td></td>
</tr>
<tr>
<td>Female</td>
<td>36</td>
<td>33</td>
<td></td>
</tr>
<tr>
<td>BMI, kg/m²</td>
<td>25.1±3.2</td>
<td>24.6±3.4</td>
<td>0.41</td>
</tr>
<tr>
<td>Smoking</td>
<td>37</td>
<td>34</td>
<td>0.69</td>
</tr>
<tr>
<td>Hypertension</td>
<td>47</td>
<td>51</td>
<td>0.52</td>
</tr>
<tr>
<td>Diabetes</td>
<td>24</td>
<td>22</td>
<td>0.61</td>
</tr>
<tr>
<td>Family history of CHD</td>
<td>39</td>
<td>24</td>
<td>0.23</td>
</tr>
<tr>
<td>Medication</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Nitric acid ester</td>
<td>60</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td>β-blocker</td>
<td>72</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td>Antithrombotic</td>
<td>63</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td>Statins</td>
<td>87</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td>Other</td>
<td>28</td>
<td>NA</td>
<td>NA</td>
</tr>
</tbody>
</table>

For smoking, hypertension and diabetes categories: The individuals who smoked at the time of the study or previously in their lifetime and who had a history of smoking but quit smoking for less than a year were all considered smoking; individuals who had hypertension or diabetes at the time of the study or prior to the study were categorized as hypertension or diabetes. The number of individuals in each group that were smokers or had hypertension or diabetes are presented, respectively. Within the medication category, the ‘other’ group included calcium channel blockers and renin-angiotensin system inhibitors. Certain patients were treated with two drugs and others were treated with interventional therapy and medication. BMI, body mass index; CHD, coronary heart disease; NA, not applicable.
change the partial structure of the protein and be one of the major functional sites in the active region of the eNOS enzyme. As a result of altered structure of the active region of eNOS, the activity of enzyme may subsequently be altered by affecting its ability to bind to substrates, thus affecting biological functions.

**Signalizing pathway associated with eNOS and CHD.** KEGG pathway analysis demonstrated that eNOS is involved in the processes of atherosclerosis and arginine and proline metabolism. In addition, eNOS was demonstrated to be a key enzyme involved in the conversion of L-arginine to NO in endothelial cells (22); eNOS catalyzes the generation of NO, which activates guanylate cyclase in vascular smooth cells to increase cyclic (c)GMP levels, which subsequently relaxes the vascular smooth muscle.

![Figure 2. Tertiary structure diagram for the partial structure of the G894T endothelial nitric oxide synthase polymorphism, which obtained using QUARK and the Cn3D viewer software. (A) Protein structure when the 298th amino acid was glutamic acid. The modeling sequence was 'lplllqapd-dppelfllppe'. (B) Protein structure when the 298th amino acid was aspartic acid. The modeling sequence was 'lplllqapdeppelfllppe'.]
smooth muscle and leads to vasodilation to regulate the blood pressure, and consequently prevents cardiovascular diseases such as atherosclerosis (23). Therefore, it may be hypothesized that, if the partial structure of eNOS is altered due to the eNOS G894T polymorphism, the activity of the enzyme may be reduced. Upon reduced enzymatic activity, the binding of L-arginine and eNOS may be reduced, which would subsequently reduce the generation of NO and lead to improper regulation of vasodilatation and vasoconstriction, with altered blood pressure and blood flow, which may reduce superoxide clearance (24). As a result, coronary vascular atherosclerosis and thrombi may develop, which increase the risk of CHD (Fig. 4). Therefore, alterations in the activity of eNOS, levels of NO, activity of guanylate cyclase and levels of cGMP in individuals with the different genotypes for this polymorphism should be determined in further studies to confirm the above hypothesis, which may aid the development of novel drugs for the treatment of CHD at a molecular level.

Discussion

CHD may be caused by various factors, including genetic and environmental factors (25). At present, CHD incidence is increasing within the Chinese population and the number of young patients with CHD is increasing annually (26). Studies have demonstrated that during the process of atherosclerosis, eNOS gene polymorphisms may affect the activities of the enzyme and be associated with the incidence of CHD (27,28). However, to the best of our knowledge, the association between eNOS gene polymorphisms and CHD in young people has not been previously clarified. The present study compared the distribution of genotypes and allele frequency of the eNOS G894T gene polymorphism between CHD and control groups, analyzed alterations in the protein structure by bioinformatics and further discussed the association between the eNOS G894T polymorphism and the risk of CHD in young individuals, in addition to the potential mechanisms.

The association between the eNOS G894T gene polymorphism and the risk of premature CHD has been reported by various studies globally, however, results obtained in the same region and for similar ethnic groups were different (15,29,30). A study by Colombo et al. (31) included 315 Italians with 201 CHD cases and 114 controls, and the results demonstrated that the incidence and severity of CHD was associated with the G894T polymorphism. By contrast, a study on an Italian population by Rossi et al. (32) indicated that CHD was independent of the eNOS G894T polymorphism. In China, He et al. (33)
reported that differences in the T allele frequency of the G894T polymorphism between the CHD and control groups were statistically significant (P<0.05) in a study where participants were from Henan province. However, Liang et al (34) demonstrated that the genotype TT was significantly different in the CHD and control groups (OR, 8.50, P<0.05), while differences in the T allele frequency of the G894T polymorphism between the CHD and control groups was not statistically significant. The results of the present study indicated that the differences in the distribution of genotype and allele frequency of the eNOS G894T gene polymorphism between the young CHD and control groups were statistically significant, which indicated that the incidence of CHD in young people may be associated with the eNOS G894T gene polymorphism. The risk of CHD associated with the GT genotype was 1.867 times the risk associated with the GG genotype, while the risk for TT and GT + TT genotypes was 2.075 and 1.889 times the risk for the GG genotype, respectively. In addition, the risk of CHD associated with the eNOS 894T allele was 1.749 times the risk associated with the eNOS 894G allele (P<0.05). These results indicated that individuals with the T allele of the G894T polymorphism may be at a higher risk of CHD. The result of structure modeling based on the amino acid sequence demonstrated that the protein structure of the G894T polymorphism was different prior to and following mutation, which led to partial structural alterations in the active structural domain of eNOS being altered from a random coil to an α-helix. Therefore, it was hypothesized that this altered structure may influence the activity of eNOS and the binding of eNOS with L-arginine, consequently decreasing the generation of NO. Low NO levels restrict vasodilatation, reduce blood flow, affect the clearing of superoxide in the blood, promote blood platelet adherence and reduces the oxidation of low-density lipoprotein cholesterol. Therefore, reduced NO levels accelerates the formation of coronary atherosclerosis and thrombi, consequently increasing the risk of CHD (35). Although the observed structural difference would be present in all individuals that possess the 894T allele, and not just young individuals with this allele, 894T may be an important risk factor for early-onset CHD. Therefore, it may be more obvious in young patients with CHD that the eNOS G894T polymorphism is associated with the development of CHD, compared with older patients. In conclusion, the occurrence and development of CHD in young people may be associated with the eNOS G894T gene polymorphism. The potential mechanism may be that the 894G→T mutation, which causes the 289th amino acid Glu→Asp missense mutation in the protein sequence, leads to alterations in the functional structure domain of partial structures in eNOS, influencing the activity of eNOS and the binding of substrates to eNOS and reducing the generation of NO, consequently leading to the incidence of CHD. However, there certain limitations are associated with the present study. Firstly, only one polymorphism site was analyzed in the current study, and the interaction between G894T and other polymorphism sites in eNOS has not been investigated. Previous studies have indicated that additional polymorphism sites in eNOS, including rs2070744, T786C and eNOS4a/4b 27 bp variable number tandem repeat, may also be associated with CHD (22,36,37); therefore, whether the G894T polymorphism may lead to CHD via combined action with other polymorphism sites in eNOS requires further investigation. Additionally, the mechanism was analyzed by a bioinformatics tool and, although structural alterations were identified, the effect of these structural alterations on eNOS activity requires verification, which may illustrate the association between the G894T polymorphism and the risk of CHD more clearly. Furthermore, although the risk of CHD was analyzed in the current study, the association between the G894T and the efficacy for drugs of the treatment of CHD was not determined. Therefore, investigation into the association between the eNOS polymorphism and the efficacy for drugs of the treatment of CHD is required to determine whether the polymorphism affects drug sensitivity or multi-drug resistance may be of clinical importance for the treatment of CHD.

References