High risk of essential hypertension in males with intron 4 VNTR polymorphism of eNOS gene

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In this study 250 patients with essential hypertension were investigated in comparison to 218 normotensives for association with epidemiological parameters. Of these DNA samples from 176 patients and 168 controls were analyzed for intron 4 27bp repeat polymorphism of eNOS gene. The study revealed significantly high risk of essential hypertension for individuals who were obese, with a positive family history and with non-vegetarian food habits. Though the intron 4b/a polymorphism of eNOS gene did not reveal any association with essential hypertension in general, males with a/a genotype of the polymorphism did show significantly high risk for developing hypertension.

Key words: Essential hypertension, endothelial NOS gene, polymorphism

Introduction

Hypertension is a multi-factor disease involving interaction of both environment and genetic components. It is a major risk factor for Coronary Artery Disease (CAD) which is associated with high mortality rate. Clinical and experimental studies have suggested involvement of several genes in the causation of hypertension. Alteration in Nitric Oxide (NO) metabolism is considered a major contributing factor. NO, a powerful endogenous vasodilator, is synthesized by endothelial Nitric Oxide Synthase gene (eNOS) and inhibits the adhesion, and recruitment of platelets, vascular smooth muscle cell migration and its growth. It also limits the oxidation of atherogenic low density lipoproteins.

The constitutive endothelial NOS (eNOS) expressed in endothelium is encoded by a gene on chromosome 7q35-36 position. It comprises of 26 exons that spans 21 kilo bases encoding an mRNA of 4052 nucleotides. Several polymorphisms of eNOS gene are found to be associated with increased risk for CVD. Of these 894 G greater than T variant in exon 7 is reported to be associated with CAD while 786 T greater than C polymorphism has been associated with Hypertension and with coronary spasm. It is considered a risk factor for CAD in Caucasians.

A 27 bp VNTR located in intron 4 of eNOS gene was proven to be of equal interest. Wang et al. reported a significant association of this intron with CAD. The study identified two alleles in intron 4, the larger allele with five tandem repeat units of 27 bps with first three having A and the last two G at 19th position of the repeat unit [GAAGTCTAGACCTGCTGC(A/G)GGGGTGAG] and the smaller allele with only four repeats, in which the first two repeats had A and the last two had G at the 19th position. The authors have designated these alleles as eNOS 4a for shorter allele with four repeats and eNOS 4b for larger allele with five repeats. The study also showed that homozygosity for allele 4a had higher risk for CAD among smokers.

Other studies on eNOS intron 4 polymorphism showed positive association with renal disease and essential hypertension among Japanese and stroke among Chinese. However, studies from Taiwan did not reveal any association with premature CAD. The discrepancy in these studies on the association of eNOS intron 4b/a VNTR polymorphism with Essential Hypertension may be related to ethnic diversity. Hence we investigated this polymorphism in Indian patients with Essential Hypertension to find the extent of risk, caused by this gene, if any.

Key words: Essential hypertension, endothelial NOS gene, polymorphism

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Original Article

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Materials and Methods

Study subjects

The study population included 250 individuals with essential hypertension diagnosed according to World Health Organization (WHO) criteria. Subjects included in the study comprised of those who were diagnosed as primary hypertensives by physicians (based on clinical and other investigations) and those who were already on antihypertensive drugs at the time of study. The patients were recruited at Gandhi Medical College and Hospital, Hyderabad, India. It is well equipped with diagnostic facilities. Patients of all socio-economic groups visit this hospital. Subjects diagnosed with secondary hypertension arising due to CAD, renal failure and other associated conditions were excluded from the study. The data generated was analyzed in comparison to that found in 218 normotensive healthy subjects that formed a control group.

Detailed information relating to the age, sex, BMI, Family history, consanguinity, diet and habits like alcohol consumption and smoking were collected from both patients and controls. Family information in terms of a four generation pedigrees was also obtained. Of the cases and controls registered for the study, 5 ml of venous blood was collected in EDTA vaccutainers from 176 cases and 168 controls. Informed consent was obtained from each individual for their participation in the study which was approved by the Ethical Committee of our institution.

Analysis of VNTR Polymorphism of eNOS

Genomic DNA was extracted from peripheral blood leukocytes by non-enzymatic method. Genotypes for eNOS polymorphism were determined by polyacrylamide gel electrophoresis after PCR amplification (Biometra, Germany) of the target region. The primers used for amplification were forward 5’- AGGCCCATATGGTAGTGCCTT -3’ and the reverse 5’- TCTCTTAGTGCTGTGGTAC -3’ that flank the region of the 27 bp repeat in intron 4 of eNOS gene. Each reaction mixture was heated to initial denaturation of 94°C for 4 minutes with 35 cycles of denaturation at 94°C for 1 minute, annealing at 56°C for one minute, extension at 72°C for two minutes and a final extension at 74°C for seven minutes.

The PCR products were run on eight per cent PAGE gels (Biotech, Indigenous), and the fragments separated were visualized by ethidium bromide staining under UV trans-illumination. PCR analysis of genomic DNA generated fragments of 393 bps corresponding to 4a/a homozygotes, 420 bp to 4b/b homozygotes and 393 and 420 bp 4b/a heterozygote [Figure 1].

The data was analyzed using descriptive statistics for epidemiological parameters and test of significance and odds ratio estimation to evaluate the risk of VNTR genotypes at eNOS locus in terms of association with hypertension.

Results

Table 1 shows the base line features found in the patient and control groups. The mean blood pressure recorded in hypertensive patients was 152.83 plus/minus 23.28 for SBP and 95.55 plus/minus 14.12 for DBP. The males and females occurred with equal incidence in the present hypertensive patients with the mean age group of 52.73 plus/minus 11.52. The frequency of case with positive family history was significantly higher in hypertensives (51.2%) as compared to controls (48.2%; \( \chi^2 = 8.105, P = 0.004 \)) suggesting the presence of strong genetic component underlying hypertension. The frequency of obese individuals was significantly higher among patients (13.2%) as compared to controls (2.75%; \( \chi^2 = 16.64, P = 0.00005 \)). Similarly, the frequency of non-vegetarians...
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was found to be higher in the patient group (85.6%) as in comparison to controls (77.06%; \( \chi^2 = 5.657, P = 0.017 \)). There was no significant deviation seen among smokers and alcoholics between patient group and control groups.

The analysis of polymorphism at eNOS intron-4 showed an incidence of 65.9% of b/b, 28.4% of b/a and 5.6% of a/a genotypes in cases and 69% of b/b, 27.9% of b/a, 2.9% of a/a among controls [Table 2]. The gene and genotypic frequencies in hypertensives and controls were in agreement with Hardy-Weinberg Equilibrium. The distribution of eNOS genotypes when tested didn’t show significant deviation between patients and controls [\( \chi^2 = 1.57 \) with 2 df, \( P = 0.455 \)]. However significant deviation was observed between male patients and male controls where b/a heterozygotes (31.4%) and a/a homozygotes (9.3%) were higher in the patient group as compared to controls [b/a 26.0% and a/a 2.0%; \( \chi^2 = 6.18, P = 0.045 \)].

There was no significant deviation seen among smokers and alcoholics between patient group and control groups.

A significant deviation in distribution of genotypes was also observed between males and females within hypertensives group where in the frequency of b/b genotypes was higher in females (72.2%) as compared to males [59.3%; \( \chi^2 = 5.52, P = 0.06 \)]. To compute the odd’s ratios the genotypic frequencies were pooled in different combinations to estimate the risk for the condition. When distribution of b/b genotype was compared with pooled b/a and a/a genotypes, a significant deviation was seen in males [\( \chi^2 = 3.32, P = 0.068, \text{OR} = 0.566, \text{with 0.307-1.066 CI at 10\% level} \)]. In compliance to this when a/a genotype was compared to pooled b/b and b/a genotypes, a significant result was obtained [\( \chi^2 = 4.846, P = 0.027, \text{OR} = 5.025, \text{with 1.307-24.34 CI} \)]. Further when the frequencies of 4b/a alleles were tested, significant OR was obtained in males supporting high risk of hypertension for males carrying allele ‘a’ [\( \chi^2 = 2.93, P = 0.086, \text{OR} = 0.529, \text{with 0.254-1.1035 CI} \)].

**Discussion**

It was demonstrated that 27bp repeat in the eNOS gene could bind nuclear proteins as an enhancer/repressor to promote/suppress the transcription efficiency. Functional significance of this polymorphism was also identified in

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**Table 1: Distribution of epidemiological parameters in hypertensive and normotensive groups**

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Hypertensives</th>
<th>Controls</th>
<th>( \chi^2 )</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>( n )</td>
<td>%</td>
<td>( n )</td>
</tr>
<tr>
<td>Total</td>
<td>250</td>
<td>218</td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>125</td>
<td>50.0</td>
<td>131</td>
</tr>
<tr>
<td>Female</td>
<td>125</td>
<td>50.0</td>
<td>87</td>
</tr>
<tr>
<td>Obese</td>
<td>33</td>
<td>13.2</td>
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</tr>
<tr>
<td>Non-obese</td>
<td>217</td>
<td>86.8</td>
<td>160</td>
</tr>
<tr>
<td>Smokers</td>
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<td>22.0</td>
<td>82</td>
</tr>
<tr>
<td>Non-smokers</td>
<td>195</td>
<td>78.0</td>
<td>136</td>
</tr>
<tr>
<td>Alcoholic</td>
<td>95</td>
<td>38.0</td>
<td>83</td>
</tr>
<tr>
<td>Non-alcoholic</td>
<td>155</td>
<td>62.0</td>
<td>135</td>
</tr>
<tr>
<td>Familial</td>
<td>128</td>
<td>51.2</td>
<td>50</td>
</tr>
<tr>
<td>Non-familial</td>
<td>122</td>
<td>48.2</td>
<td>168</td>
</tr>
<tr>
<td>Veg</td>
<td>36</td>
<td>14.4</td>
<td>6</td>
</tr>
<tr>
<td>Non-veg</td>
<td>214</td>
<td>85.6</td>
<td>212</td>
</tr>
<tr>
<td>SBP</td>
<td>152.83 ± 23.28</td>
<td></td>
<td></td>
</tr>
<tr>
<td>DBP</td>
<td>95.55 ± 14.12</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

\( \chi^2 = 6.18, *P = 0.045 \)

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**Table 2: Distribution of eNOS intron 4 b/a genotypes with reference to epidemiological parameters in hypertensives and normotensives**

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Hypertensives</th>
<th>Controls</th>
<th>( \chi^2 )</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>( n )</td>
<td>%</td>
<td>( n )</td>
</tr>
<tr>
<td>Total</td>
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<td>65.9</td>
<td>50</td>
</tr>
<tr>
<td>Male</td>
<td>51</td>
<td>59.3</td>
<td>27</td>
</tr>
<tr>
<td>Female</td>
<td>65</td>
<td>72.2</td>
<td>23</td>
</tr>
<tr>
<td>Obese</td>
<td>17</td>
<td>65.4</td>
<td>8</td>
</tr>
<tr>
<td>Non-obese</td>
<td>99</td>
<td>66.0</td>
<td>42</td>
</tr>
<tr>
<td>Smokers</td>
<td>21</td>
<td>55.2</td>
<td>15</td>
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<tr>
<td>Non-smokers</td>
<td>96</td>
<td>69.02</td>
<td>35</td>
</tr>
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<td>Alcoholic</td>
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<td>62.5</td>
<td>20</td>
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<tr>
<td>Non-alcoholic</td>
<td>76</td>
<td>67.8</td>
<td>30</td>
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<tr>
<td>Familial</td>
<td>60</td>
<td>62.5</td>
<td>31</td>
</tr>
<tr>
<td>Non-familial</td>
<td>55</td>
<td>68.7</td>
<td>20</td>
</tr>
<tr>
<td></td>
<td>116</td>
<td>69.0</td>
<td>47</td>
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</tbody>
</table>

\( \chi^2 = 6.18, *P = 0.045 \)
cases with endothelial dysfunction. In the present study the frequency of 4a allele was found to be approximately 0.16, which is little higher than that found in other populations viz., Iranian (0.1), Japanese (0.1 to 0.13) and Turkish (0.14) but is slightly lower than in Australian (0.17) and African Americans (0.26). The differences in the ethnic origin and sample sizes studied might have caused the differences in the distribution of eNOS intron 4a polymorphism studied in these populations. With reference to diseases Wang et al. showed a significant association of eNOS polymorphism with CAD in smokers but not with hypertension. Uwabo et al. have shown a significant association of hypertension with a/a homozygosity among the Japanese. They also suggested that this VNTR polymorphism may be in linkage disequilibrium with other genes related to essential hypertension. The report of Ichihara et al. and Salmi et al. have shown a positive association of eNOS 4b/a polymorphism with CAD in Japanese and Iranian populations. Absence of such association was reported in German and Taiwanese populations.

Though this study did not reveal significant association of eNOS intron4 polymorphism with essential hypertension in general, males with carriers for allele ‘a’ did show a significant risk for the condition which was five times higher compared to other genotypes. Further, individuals who were obese, with positive family history, and non-vegetarian food habits were also at risk as evidenced by the high occurrence of hypertensives in these groups. Contribution of these risk factors has to be evaluated with large samples and cohorts of hypertensives.

Acknowledgment

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References

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Announcement

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41st American Cytogenetics Conference

Conference dates: May 13-16, 2010
Venue: Crowne Plaza “Brock Plaza” Hotel Falls Avenue, Niagara Falls, Ontario, Canada. Directly opposite to the “Rainbow Bridge”
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Identify: American Cytogenetics Conference for Conference rates
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Shuttle service available upon request - 24 hr prior to arrival from: Toronto International & Buffalo Niagara International Airports

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Abstracts: 250-300 words (Times New Roman). Follow ASHG abstract style

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For more information – Contact: Avirachan T. Tharapel, PhD – President, American Cytogenetics Conference atharapel@utmem.edu

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Source of Support: UGC, India.
Conflict of Interest: None declared.

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- Example of a correct style
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- Add up to a maximum of 15 references at a time.
- If the reference is correct for its bibliographic elements and punctuations, it will be shown as CORRECT and a link to the correct article in PubMed will be given.
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