Frequencies and Distribution of APOE Gene Polymorphisms and Its Association With Lipid Parameters in the Senegalese Population

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Abstract

Background

Apolipoprotein E is a multifunctional protein that plays an important role in lipid metabolism. It is encoded by the APOE gene. However, APOE gene polymorphism has not been very well studied in the Senegalese population. Therefore, we studied allele frequencies, genotype distributions, and the relationship between APOE gene polymorphisms and lipid parameters in the Senegalese women population.

Methodology

This study included 110 healthy women aged 35-72 years. The mean age was 49.8 ± 8.1 years. For all subjects, lipid parameters were analyzed from fasting serum, and APOE genotypes were identified by PCR-RFLP (polymerase chain reaction-restriction fragment length polymorphism) based analysis.

Results

Variations in the frequencies and distribution of the APOE alleles and genotypes were observed (ε3: 47.3%; ε2: 45.2%; ε4: 9.6%; and ε2/ε3: 70%; ε2/ε4: 16.4%; ε3/ε3: 10.9%; ε2/ε4: 2.7%). Compared to the ε3/ε3 genotype carriers, carriers of the ε3/ε4 genotype had a significantly higher rate of total cholesterol (p=0.03) and no high-density lipoprotein-cholesterol (p=0.02). Univariate analysis showed that the APOE ε4 allele increases the low-density lipoprotein-cholesterol rate (OR=3.06; 95% CI: 1.16-8.22; p=0.02).

Conclusion

Our study has shown a difference in APOE allele frequencies and genotype distributions with a total absence of ε2ε2 and ε4ε4 genotypes in a sample of Senegalese women. We also found that APOE gene polymorphism might play a role in plasma lipid levels.

Categories: Genetics, Epidemiology/Public Health, Nutrition

Keywords: apolipoprotein e, senegalese, lipids, frequencies, polymorphism

Introduction

Apolipoprotein E (ApoE) is a polymorphic and multifunctional protein that is mainly synthesized by the liver. It is an essential apolipoprotein in plasma [1] and plays a vital role in the transport, metabolism, and digestion of lipoproteins [2-4]. ApoE promotes efficient clearance of circulating lipoproteins and participates in the cellular efflux of cholesterol [5]. ApoE plays an important role in regulating lipoprotein metabolism by regulating the binding of these lipoproteins to specific receptors. It can bind with different molecules, such as low-density lipoprotein-cholesterol (LDL-C), high-density lipoprotein-cholesterol (HDL-C), very low-density lipoprotein-cholesterol (VLDL-C), and chylomicron to participate in the transformation and metabolism of lipoprotein.

Human ApoE is encoded by the APOE gene (OMIM: 107741), which is located on chromosome 19q13.32 [6,7]. The APOE gene is polymorphic in nature and possesses three alleles, namely, ε2, ε3, and ε4, which can be found in six genotypic combinations, homozygous (ε2/ε2, ε3/ε3, ε4/ε4) or heterozygous (ε2/ε3, ε2/ε4, ε3/ε4) [8,9].

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The APOE gene regulates ApoE plasma concentration and its binding capability. The three alleles (ε2, ε3, ε4) of the APOE gene are respectively responsible for the production of corresponding apolipoprotein E2, apolipoprotein E3, and apolipoprotein E4 plasma isoproteins [10]. Different alleles lead to structural variations in ApoE protein (E2, E3, E4), influence its functions, and contribute to the variation in lipoprotein concentration, including receptor-binding capacity and lipid metabolism.

Therefore, the ε2, ε3, and ε4 genetic variants and their corresponding protein variations (E2, E3, and E4) have been linked to differential risks of dyslipidemia [11]. Dyslipidemia further confers as a risk factor of atherosclerosis, cerebral infarction, diabetes, and hypertension. Preventing dyslipidemia plays an important role in reducing morbi-mortality rate in the world [12]. At present, there are limited studies on the Senegalese populations. Hence, in the present study, we examined the allelic and genotypic frequencies of the APOE gene and its role in the lipid parameters in Senegalese women subjects firstly.

Materials And Methods

Study participants and protocol

A total of 110 healthy subjects were included in this study. According to their ethnicity, there were 104 Wolofs, 49 Peulhs, 44 Toucouleurs, 15 Diolas, 15 Lebous, 8 Bambaras, 3 Soces, and 19 other ethnicities. The age of the participants in the study was between 35 and 72 years, and the mean age was 49.79 ± 8.10 years. They were recruited at the physiology of the Cheikh Anta Diop University (UCAD, Dakar, Senegal). The subjects were HIV, HBV, and HCV negative.

Inclusion criteria were as follows: > 18 years of age, females, and having not taken lipid-regulating drugs before or having stopped taking the drug for at least three months. Subjects were excluded if they were diagnosed with a disease that affected blood lipid level. Pregnant and breastfeeding women were also excluded.

All procedures were conducted in accordance with the standards of the Declaration of Helsinki. It was reviewed and approved by the Ethics Committee of UCAD (Reference: Protocole 027512018/CERruCAD). All study participants provided signed informed consent.

Lipid analysis

Approximately 5 mL of fasting venous blood samples (at least 12 hours of overnight fasting) was collected into vacutainers from every participant and stored at -80°C in a deep freezer until further analysis.

All biochemistry parameters were analyzed on an automated Abbott device (ARCHITECT i1000SR, Abbott Laboratories, Abbott Park, Seattle, WA, USA) according to the standard laboratory protocol. The serum samples were assayed for blood lipid profiles. On the heparin tube, we measured the following lipids: apolipoprotein A (Apo A) apolipoprotein B (Apo B), total cholesterol (TC), HDL-C, LDL-C, and triglycerides (TG). TC, HDL-C, and TG were measured by the enzymatic method. Apo A and Apo B were measured by enzyme immunology.

LDL-C level was calculated according to the Friedewald equation: LDL-C = TC − HDL-C − TG/5. No-HDL-C level was calculated by subtracting HDL-C value from the TC value: No-HDL-C = TC − HDL-C.

DNA extraction and genomic assay

DNA Extraction

Genomic DNA was extracted from peripheral blood lymphocytes using a commercial kit (REF A1125, Wizard® Genomic DNA Purification kit, Promega Corporation, Madison, WI, USA), as per the manufacturer’s instructions. The DNA concentration was quantified using a NanoDrop 2000™ spectrophotometer (ThermoFisher Scientific, Waltham, MA, USA).

PCR-RFLP based analysis

APOE gene was amplified by polymerase chain reaction (PCR) on ThermoCycler (T-personal, Biometra, Jena, Germany). The PCR conditions included initial denaturation phase at 95°C for 5 minutes followed by 35 cycles of denaturation at 95°C for 30 seconds, annealing at 70°C for 20 seconds, extension phase at 72°C for 20 seconds, and the final elongation at 72°C for 10 minutes. PCR amplification was visually confirmed on 1.5% agarose gel electrophoresis.

After confirmation of amplification, 10 µl of each amplified DNA was mixed with each of the two restrictions enzymes (HaeII and AFLIII). The two reactions were allowed to proceed for at least 3 hours at 57°C. The resulting fragments were analyzed on a 4% agarose gel (Figure 1).
FIGURE 1: PCR-RFLP gel with AfLIII and HaeII enzyme.

ε2/ε2: one band at 227 bp for the HaeII and two bands (at 177 and 50 bp) for AfLIII. ε3/ε3: two bands (at 195 and 32 bp) for the HaeII and two bands (at 177 and 50 bp) for AfLIII. ε4/ε4: two bands (at 227 bp for AfLIII and one band at 227 bp for HaeII. ε2/ε3: three bands (at 227, 195, and 32 bp) for the HaeII and two bands (at 177 and 50 bp) for AfLIII. ε2/ε4: three bands (at 227, 195, and 32 bp) for the HaeII and three bands (at 227, 177, and 50 bp) for AfLIII. ε3/ε4: two bands (at 195 and 32 bp) for the HaeII and three bands (at 227, 177, and 50 bp) for AfLIII.

PCR-RFLP, polymerase chain reaction-restriction fragment length polymorphism

Statistical analysis

The data were expressed as mean ± SD and percentages (%). Allele frequencies were estimated by the gene counting method, and the chi-square test was used to identify departure from Hardy-Weinberg equilibrium. Strength of association between different lipid variables and APOE genotypes and alleles was estimated using the chi-square test and logistic regression. SPSS statistical software Version 23.5 (IBM Corp., Armonk, NY, USA) was used for data analysis. A p-value of ≤0.05 was considered as significant in statistical analysis.

Results

APOE alleles and genotype frequencies

We found that the ε3 allele is the predominant form while the ε2ε3 genotype is the most common form in the study population. In addition, there is a total absence for ε2ε2 and ε4ε4 (Table 1).
### Variables

<table>
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<th>Frequency of APOE alleles</th>
<th>Subjects</th>
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<tbody>
<tr>
<td>Allele n (%)</td>
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<tr>
<td>ε2</td>
<td>95 (43.18)</td>
</tr>
<tr>
<td>ε3</td>
<td>104 (47.27)</td>
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<tr>
<td>ε4</td>
<td>21 (9.55)</td>
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</tr>
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</tr>
<tr>
<td>ε2ε3</td>
<td>77 (70.00)</td>
</tr>
<tr>
<td>ε2ε2</td>
<td>/</td>
</tr>
<tr>
<td>ε2ε4</td>
<td>18 (16.36)</td>
</tr>
<tr>
<td>ε3ε4</td>
<td>3 (2.73)</td>
</tr>
<tr>
<td>ε4ε4</td>
<td>/</td>
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</tr>
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<tr>
<td>p-value</td>
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</table>

**TABLE 1: Characteristics of APOE gene polymorphisms**

VA, allelic variation

**Comparison of lipid profile between participant subgroups according to APOE gene polymorphisms**

The ε3ε4 genotype compared to the ε3ε3 genotype had a significantly higher mean level of TC and no-HDL-C (Table 2). We also found that only ε4 carriers had abnormal mean levels of total and LDL-C and that the mean TG level is normal regardless of the genotype considered.
TABLE 2: Difference in the lipid profile of the subjects according to APOE genotype

*A p-value of ≤0.05 was considered as significant in statistical analysis using the ANOVA test. The post hoc test was the least significant difference test.

Apo A, apolipoprotein A; Apo B, apolipoprotein B; HDL, high-density lipoprotein; LDL, low-density lipoprotein; TC, total cholesterol; TG, triglycerides

We also found that carriers of the ε4 allele had higher blood LDL-C levels (Table 3).

TABLE 3: Associations between lipid parameters and APOE allele (categories of lipid parameters are normal and abnormal rates of each parameter)

*A p-value of ≤0.05 was considered as significant in statistical analysis using chi-square and odds ratio tests.

Apo A, apolipoprotein A; Apo B, apolipoprotein B; HDL, high-density lipoprotein; LDL, low-density lipoprotein
Discussion

ApoE, one of the major protein components of lipoproteins, plays an essential role in the circulation and the metabolism of blood lipids [13]. It is encoded by a gene that is characterized by polymorphisms. At present, there are limited studies on APOE gene polymorphism in the Senegalese population. Correlation between allelic and genotypic frequencies of APOE polymorphisms versus the risk of dyslipidemia served as a key attraction to conduct this study.

In this study, the analysis showed the three alleles of the APOE gene, which are all present in the Senegalese population with frequency distribution as ε3(47.3%)>ε2(43.2%)>ε4(9.6%). Furthermore, it should be mentioned that the APOE genotype distribution in this study was ε2ε3(70%)>ε2ε4(16.4%)>ε3ε4(10.9%)>ε3ε3(2.7%). However, some genotypes such as ε2ε2 and ε4ε4 were totally absent in our study population of adult subjects. We estimate that the absence of the ε2ε2 and ε4ε4 genotypes, in our study population consisting of elderly subjects, would be justified by a relatively short life expectancy for individuals harboring these genotypes. In effect, ε2ε2 homozygosity can precipitate type III hyperlipoproteinemia [14] while ε4ε4 would increase plasma LDL levels and the risk for atherosclerosis [2]. This differs from the results of some studies conducted in other populations such as in Algeria [15], where all APOE genotypes were present, and in another study from Saudi Arabia, researchers noted a total absence of ε2 and consequently the absence of the corresponding genotypes ε2ε3, ε2ε4, and ε2ε2 [16].

In the study, APOE ε2ε3 is the most common genotype and ε3 is the most predominant allele. This is contrary to previous studies because the ε3 allele is generally the most predominant allele but the most common genotype was ε3ε3 [17,18], and the worldwide frequency of the ε2, ε3, and ε4 alleles is 8.4%, 77.9%, and 13.7%, respectively [19]. Authors have reported that the ε2 isoform is the least common [15], which does not support our results, although the frequencies of APOE alleles and genotypes vary considerably between different ethnicities and populations [20].

In our study, we found that individuals harboring ε4 allele has higher blood LDL-cholesterol levels (OR=3.06; 95% CI: 1.16-8.22; p=0.02). In addition, the study shows that the individuals harboring ε3ε4 genotype compared to the individuals harboring ε3ε3 genotype had significantly a higher mean levels of TC (p=0.05) and no-HDL-C (p=0.02). Our results are in line with the published studies. In the literature data, the ε4 isoform was associated with an increase in the TC and LDL-cholesterol concentrations when the ε3/ε3 homozygote carriers are used as references. Similar results have been observed in different studies [21,22]. Our study has shown a significant association between ε2 allele and the plasma TG levels (OR=1.05; 95% CI: 1.00-1.09). Several previous studies rather reported increased plasma TG levels in individuals harboring allele ε2 in healthy populations [23-25] and that TG concentration was significantly lower in the carrier of ε5 allele than in the carriers of ε2 or ε4 allele [21].

ApoE isoforms differ in their binding affinity to serum cholesterol and hence in their ability to remove dietary fats from the blood [26]. Accordingly, the total serum cholesterol levels differ between APOE alleles and genotypes [26,27]. In view of these findings, it appears that despite differences of only one or two amino acids, the structural and functional differences between the three apoE isoforms may have a profound effect on disease risk [28]. The mature form of ApoE has two structural domains separated by a hinge region. The amino-terminal domain (amino acids 1-191) contains the LDL receptor-binding region, and the carboxyl-terminal domain (amino acids 225-299) contains the lipid-binding region [2]. The structural basis of the three isoforms occurs through amino acid exchanges of amino acid residues of the polypeptide chain at positions 112 and 158 of the protein sequence where cysteine (Cys) or arginine (Arg) is present. The most common isoform, ApoE3, has a Cys at position 112 and an Arg at position 158 (Cys112, Arg158), ApoE2 has a Cys at position 112 and an Arg at position 158 (Cys112, Cys158), and ApoE4 has an Arg at positions 112 and 158 (Arg112, Arg158) [2]. In ApoE3, Arg-158 forms a salt bridge with aspartic acid-154, whereas in ApoE4, with Cys-158, the salt bridge is interrupted and aspartic acid-154 interacts with Arg-150, modifying the whole region receptor binding [2]. ApoE2 isoform shows defective binding to hepatic lipoprotein receptors [28]. Thus, the differential abilities of ApoE isoforms to bind to hepatic lipid receptors may contribute to ApoE isoform-specific effects in disease. This was confirmed by the study of Eto et al., which showed that the E2 allele and the E4 allele are associated with an increased risk of ischemic heart disease compared to the E3 allele [29].

The limits of our work are imputed to the study population in view of its small number and its exclusive composition of women. This study focused exclusively on a female population. In fact, in the literature, it is demonstrated that obesity is associated with polymorphisms of the APOE gene. In view of the variations in the prevalence of obesity according to gender, we wanted to first conduct a study in a female population and then, if possible, in an exclusively male population.

Conclusions

Our study has shown a significant difference in APOE allele frequencies and genotype distributions in the Senegalese women compared to other populations. The total absence of ε2ε2 and ε4ε4 genotypes in this Senegalese adult population requires a resumption to work on a younger cohort to better establish the results. We also found in the study that APOE gene polymorphism might play a role in determining plasma lipid levels.
References


