The effects of an intronic polymorphism in TOMM40 and APOE genotypes in sporadic inclusion body myositis

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A previous study showed that, in carriers of the apolipoprotein E (APOE) genotype ε3/ε3 or ε3/ε4, the presence of a very long (VL) polyT repeat allele in “translocase of outer mitochondrial membrane 40” (TOMM40) was less frequent in patients with sporadic inclusion body myositis (sIBM) compared with controls and associated with a later age of sIBM symptom onset, suggesting a protective effect of this haplotype. To further investigate the influence of these genetic factors in sIBM, we analyzed a large sIBM cohort of 158 cases as part of an International sIBM Genetics Study. No significant association was found between APOE or TOMM40 genotypes and the risk of developing sIBM. We found that the presence of at least 1 VL polyT repeat allele in TOMM40 was significantly associated with about 4 years later onset of sIBM symptoms. The age of onset was delayed by 5 years when the patients were also carriers of the APOE genotype ε3/ε3. In addition, males were likely to have a later age of onset than females. Therefore, the TOMM40 VL polyT repeat, although not influencing disease susceptibility, has a disease-modifying effect on sIBM, which can be enhanced by the APOE genotype ε3/ε3.

1. Introduction

Sporadic inclusion body myositis (sIBM) is known as the most common acquired myopathy among people 50 years and older (Machado et al., 2014). Given its similarities with Alzheimer’s disease such as the late age of onset and the abnormal accumulation of proteins, the apolipoprotein E (APOE, OMIM#107741) gene has been one of the most popular genes studied in sIBM (Gang et al., 2014), but no association with sIBM disease risk was confirmed.
2. Methods

A total of 158 DNA samples from biopsy-confirmed sIBM patients (66.5% males) were collected from 9 centres around the world. The control group comprised 127 individuals with no history of neuromuscular disease. Restriction fragment length polymorphism was used for APOE genotyping. Fluorescence-based fragment size analysis was performed for genotyping the polyT repeat in intron 6 of the TOMM40 gene (rs10527454). The polyT repeat alleles were classified based on the length of repeat (N) as originally established by Roses et al. (2010). Genotypic frequencies for APOE and TOMM40 polyT were compared between cases and controls using the chi-square or Fisher’s exact test. The associations between APOE genotypes, TOMM40 polyT genotypes, ethnicity, gender, and age of onset of sIBM were analyzed by linear regression analyses. For all the analyses, p value <0.05 was considered statistically significant (for details, see Supplementary Data).

3. Results

The APOE ε2/ε2 genotype was absent in both the sIBM cohort and the control group. No significant differences were found between patients and controls regarding APOE allelic and genotypic frequencies, considering neither only Caucasian patients nor the entire cohort. The distribution of TOMM40 polyT repeat lengths was similar in the sIBM group (range 8–36 repeats) and controls (11–33 repeats) (Supplementary Fig. 1). The frequency of neither TOMM40 polyT repeat genotypes nor polyT alleles differed significantly between groups (Supplementary Table 1). When the analysis was performed in the combined group of carriers of ε3/ε3 and ε3/ε4 APOE genotypes, there was still no association between the presence of VL polyT repeats and sIBM risk (Supplementary Table 2).

There was no significant association between different APOE alleles and the age of onset of sIBM. However, being a carrier of the polyT repeat, VL allele was significantly associated with a later age of onset by 3.7 years in average (95% confidence interval [CI] = 0.4, 6.9; adjusted p = 0.027). We repeated the analysis just for carriers of the APOE ε3/ε3 genotype. We found that APOE ε3–TOMM40 VL allele carriers had an even later age of onset of sIBM by 4.9 years in average (95% CI = 1.1, 8.7; adjusted p = 0.013). Similar results were found when the analysis was restricted to Caucasians (by 4.0 years for a VL carriage, 95% CI = 0.4, 7.6; p = 0.028; by 5.4 years for carriage of a VL allele and APOE ε3/ε3 genotype, 95% CI = 1.2, 9.7; p = 0.013). Although not reaching statistical significance, there was also a trend for a later age of onset (by 2.7 years) in males compared with females (Table 1).

Table 1

<table>
<thead>
<tr>
<th>Variable</th>
<th>Count</th>
<th>Age of onset (mean ± SD), y</th>
<th>Adjusted analysis&lt;sup&gt;a&lt;/sup&gt;</th>
<th>p Value&lt;sup&gt;b&lt;/sup&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ethnicity</td>
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<td>Regression coefficient (95% CI)</td>
<td></td>
</tr>
<tr>
<td>Non-Caucasian</td>
<td>16</td>
<td>56.7 ± 5.7</td>
<td>Reference</td>
<td></td>
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<tr>
<td>Caucasian</td>
<td>141</td>
<td>60.0 ± 10.0</td>
<td>2.8 (&lt;−2.3, 7.9)</td>
<td>0.28</td>
</tr>
<tr>
<td>Gender</td>
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<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>F</td>
<td>52</td>
<td>57.9 ± 10.4</td>
<td>Reference</td>
<td></td>
</tr>
<tr>
<td>M</td>
<td>105</td>
<td>60.6 ± 9.3</td>
<td>2.7 (&lt;−0.5, 5.9)</td>
<td>0.095</td>
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<td>APOE</td>
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<td></td>
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<tr>
<td>ε2/ε2&lt;sup&gt;c&lt;/sup&gt;</td>
<td>6</td>
<td>56.8 ± 5.8</td>
<td>Reference</td>
<td></td>
</tr>
<tr>
<td>ε3/ε3</td>
<td>99</td>
<td>60.2 ± 9.8</td>
<td>−2.9 (&lt;−7.7, 1.9)</td>
<td>0.23</td>
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<tr>
<td>ε2/ε3</td>
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<td>56.9 ± 10.0</td>
<td>1.6 (&lt;−2.4, 5.7)</td>
<td>0.43</td>
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<td>ε3/ε4 and ε4/ε4</td>
<td>33</td>
<td>60.4 ± 9.7</td>
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</tr>
</tbody>
</table>

TOMM40 polyT

| No VL carriage | 74 | 58.1 ± 9.7 | Reference |                     |
| VL carriage | 83 | 61.2 ± 9.6 | 3.7 (0.4, 6.9) | 0.027 |

APOE-TOMM40

| ε3/ε3 and polyT no-VL carriage | 38 | 57.3 ± 9.9 | Reference |                     |
| ε3/ε3 and polyT VL carriage | 61 | 62.0 ± 9.4 | 4.9 (1.1, 8.7) | 0.013 |

Key: APOE, apolipoprotein E; CI, confidence interval; F, female; M, male; SD, standard deviation; sIBM, sporadic inclusion body myositis; TOMM40, translocase of outer mitochondrial membrane 40; VL, very long.

<sup>a</sup> Each analysis was adjusted for gender, ethnicity, tissue, and genetic factors, except for the variable under study.
<sup>b</sup> p value < 0.05 was considered statistically significant (marked in bold).
<sup>c</sup> ε2/ε4 was not included in the regression analysis for APOE alleles and the age of onset.

4. Discussion

This is the largest cohort where the influence of the APOE and TOMM40 genes in sIBM disease risk and features has been investigated. Concerning APOE, our findings confirmed that the APOE ε4 allele is not a susceptibility factor for developing sIBM, which is consistent with the previous studies (Needham et al., 2008). APOE alleles were also not significantly associated with the age of onset of the disease. In addition, our findings did not replicate a previously reported association between APOE–TOMM40 and risk of developing sIBM (Mastaglia et al., 2013). However, we observed that carriage of a VL repeat allele was significantly associated with a later age of onset of symptoms. This effect was even more pronounced among those also with the APOE ε3/ε3 genotype. This suggests that the TOMM40 VL polyT repeat has a disease-modifying effect on sIBM by delaying the onset of symptoms, and the APOE ε3/ε3 genotype enhances this effect. Although the association between APOE and TOMM40 and sIBM risk was not confirmed in our study, the finding of an association between the TOMM40 VL polyT repeat and a later age of onset of sIBM may justify further gene expression studies in the future. In addition, there might be other variants within the APOE–TOMM40 locus involved in the susceptibility to sIBM. Furthermore, identification of new genes and variants is crucial to improve our understanding of this complex disease.

Disclosure statement

The authors have no conflicts of interest to disclose.

Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at http://dx.doi.org/10.1016/j.neurobiolaging.2014.12.039.

(Needham et al., 2008). Recently, the “translocase of outer mitochondrial membrane 40” (TOMM40, OMIM#608061) gene, which is adjacent to and in strong linkage disequilibrium with the APOE locus on chromosome 19, has been studied in 90 Caucasian patients with sIBM (Mastaglia et al., 2013). That study reported that among carriers of the APOE genotype ε3/ε3 or ε3/ε4, carriage of a very long (VL) polyT repeat in the intron 6 of TOMM40 (rs10527454) was less frequent in sIBM compared with controls and was also associated with a later age of onset of sIBM symptoms (Mastaglia et al., 2013). To further investigate the previously reported association between APOE–TOMM40 and sIBM, we genotyped APOE and the polyT repeat polymorphism in TOMM40 in a large sIBM cohort of 158 affected cases from the International IBM Genetics Consortium and investigated their association with disease risk and the age of onset of sIBM.
References


