No Association between Single Nucleotide Polymorphisms (SNPs) of the Interferon-Induced Transmembrane Protein 3 (IFITM3) Gene and the Susceptibility of Alzheimer’s Disease (AD)

Sae-Young Won 1,2,*, Yong-Chan Kim 1,2,† and Byung-Hoon Jeong 1,2,*

Abstract: Background and Objectives: Alzheimer’s disease (AD) is the most common progressive neurodegenerative disorder, characterized by the accumulation of amyloid-beta (Aβ) in the brain. A recent study reported that the interferon-induced transmembrane protein 3 (IFITM3) protein plays a pivotal role in Aβ processing by the γ-secretase complex. Since several single nucleotide polymorphisms (SNPs) of the IFITM3 gene are related to the function and expression levels of the IFITM3 gene, the relationship between genetic polymorphisms in the IFITM3 gene and susceptibility to AD needs to be investigated. Materials and Methods: We investigated the genotype and allele frequencies of IFITM3 polymorphisms in 177 AD patients and 233 matched healthy controls by amplicon sequencing. In addition, we compared the genotype, allele and haplotype frequencies between AD patients and matched controls and performed an association analysis. Results: There were no significant differences in the genotype, allele or haplotype frequency distributions of the IFITM3 polymorphisms between AD patients and matched controls. Conclusions: To the best of our knowledge, this is the first case-control association study of the IFITM3 gene in AD.

Keywords: Alzheimer’s disease; AD; IFITM3; SNP; polymorphism

1. Introduction

Alzheimer’s disease (AD) is the most common progressive neurodegenerative disease among the several types of dementia [1–4]. AD is characterized by the accumulation of amyloid-beta-peptide (Aβ) derived from amyloid precursor protein (APP) in brain lesions. Aβ is produced by cleavage of the APP protein by the β-secretase encoded by the beta-site amyloid precursor protein cleaving enzyme 1 (BASE1) gene and γ-secretase, a complex that is comprised of a total of 4 high molecular weight subunits, including presenilin (PS), presenilin enhancer 2 (PEN-2), nicastrin and anterior pharynx defective 1 (APH-1) [5,6]. Several genetic variations of these genes affect the production of Aβ via the processing of APP protein. Thus, previous studies have reported that genetic variations in the APP, presenilin-1 (PSEN1) and presenilin-2 (PSEN2) genes are related to susceptibility to AD. In the APP gene, the genotype and allele frequencies of the -877T > C and -955A > G single nucleotide polymorphisms (SNPs) located in the promoter region were significantly associated with susceptibility to AD via transcription efficiency [7]. In addition, Q665D, K670M, N671L A692G, E693G, I716V and V717I germline mutations of the APP gene affect the onset of AD [8–10]. In the PSEN1 and PSEN2 genes, over 30 genetic variants are associated with elevated levels of Aβ40 and Aβ42 [8–13]. Based on this knowledge, several drugs targeting the APP and γ-secretase proteins have been introduced. However, there was no clear effect for any of the drugs tried [1,4]. Thus, it is necessary to discover specific targets to develop precise therapeutic agents for AD.
Interferon-induced transmembrane protein 3 (IFITM3) is a member of the innate immune system that is expressed primarily in the late endosome or lysosome and limits the entry of enveloped viruses [14–20]. In addition, a recent study have reported that upregulation of the IFITM3 protein affects γ-secretase, leading to increased Aβ production, and plays a pivotal role in the onset of AD [21]. In previous studies, the rs12252, rs34481144 and rs6598045 SNPs of the IFITM3 gene were found to be involved in the gene expression level and function of the IFITM3 protein. The rs12252 C allele is related to cleavage of the 21 amino acids at the N-terminus of the IFITM3 protein and produces an isof orm of the IFITM3 protein. The truncated IFITM3 protein induced by the rs12252 C allele has been shown to decrease the inhibition capacity of viral replication [16,22]. The A allele of the rs34481144 SNP inhibits the expression of the IFITM3 gene by increasing the CCCTC-binding factor (CTCF) binding ability at the promoter site of the IFITM3 gene [17]. The recent study has reported that rs12252 and 34481144 SNPs showed no correlation with the susceptibility of AD and the expression level of the IFITM3 protein. However, this study has performed association analysis with a relatively small sample size (control, n = 9; AD, n = 18) [21], and has not investigated rs6598045 SNP that modulates promoter activity [18]. Since several SNPs of the IFITM3 gene are related to the function and expression level of the IFITM3 gene and the IFITM3 protein plays a pivotal role in the processing of Aβ, further investigation of the relationship between genetic polymorphisms in IFITM3 and susceptibility to AD is necessary.

In the present study, we investigated genotype, allele and haplotype frequencies of the IFITM3 polymorphisms of 177 AD patients and 233 matched controls by amplicon sequencing. To find an association between IFITM3 polymorphisms and susceptibility to AD, we compared genotype, allele, and haplotype frequencies of IFITM3 polymorphisms between the two groups and performed association analysis.

2. Materials and Methods

2.1. Subjects

A total of blood samples from 177 AD patients were provided by the Pusan National University Hospital Biobank, a member of the Korea Biobank Network. A total of blood samples from 233 healthy Korean subjects were obtained from the Korea Biobank Network at the Centers for Disease Control and Prevention. The sample size used in this study may be enough to identify rare polymorphisms, including below 1% genotype frequency [22]. In addition, the sample size of healthy Korean subjects used in this study can also represent the total population of Korea with a 95% confidence level and a confidence interval of 6.42. The sample size of AD patients used in this study can also represent the total Korean AD patients with a 95% confidence level and a confidence interval of 7.36. All experimental procedures were approved according to the guidelines of the institutional review board of Jeonbuk National University and the 1964 Helsinki declaration and its later amendments or comparable ethical standards (approval number: JBNU 2019-03-009). Information for all samples was anonymized prior to investigation.

2.2. Genomic DNA Extraction

Genomic DNA was extracted from 200 µL of blood using the BeadTM Genomic DNA Prep Kit (Biofact, Daejeon, Korea) according to the manufacturer’s instructions.

2.3. Polymerase Chain Reaction (PCR)

The human IFITM3 gene was amplified using PCR with the gene-specific forward primer: 5'-CAGGGGAAGTCTCCAGGACC-3' and reverse primer: 5'-CCAAGCCACACACACACA-3'. The PCR mixture consisted of 1 µL of genomic DNA, 10 pmol of each primer, 2.5 µL of 10× Taq DNA polymerase buffer, 0.5 µL of a 0.2 µM dNTP mixture, 5 µL of 5× Band Helper, and 0.25 µL Taq DNA polymerase. The PCR conditions followed the manufacturer’s recommendations. The annealing temperature of the IFITM3 gene primers was 68 °C.
2.4. Amplicon Sequencing and Genotyping

All PCR products were purified using a FavorPrep GEL/PCR purification Mini Kit (FAVORGEN, Pingtung City, Taiwan) and sequenced on an ABI 3730 sequencer (ABI, Foster City, CA, USA). Sequencing results were visualized using Finch TV software (Geospiza Inc., Seattle, WA, USA), and genotyping was performed.

2.5. Statistical Analysis

The genotype and allele frequencies of the IFITM3 gene were compared between the AD patients and healthy controls by the chi-square \((\chi^2)\) test using SAS 9.4 software. Analysis of the Hardy-Weinberg equilibrium (HWE) test and haplotypes were performed using Haploview Version 4.2 (Broad Institute, Cambridge, MA, USA).

3. Results

3.1. Subject Description

A total of 177 Korean AD patients and 233 healthy Koreans without dementia were included in the association analysis. Detailed information on the study population is given in Table 1. The 177 AD patients included 128 females and 49 males. The 233 healthy individuals included 103 females and 130 males. The mean age of AD patients at diagnosis was 73.34 ± 7 years, and the mean age of healthy individuals at sample collection was 63.75 ± 9.74 years.

Table 1. Detailed information on the study population.

| Characteristics | Cases | Controls |
|-----------------|-------|----------|
| Number          | 177   | 233      |
| Age             | 73.19 ± 7.20 | 63.75 ± 9.74 |
| Sex (n, %)      | Male 49 | 103 |
|                 | Female 128 | 129 |
| APOE E2/E3      | 15 (8.5) | 16 (7.1) |
| E2/E4           | 7 (4) | 1 (0.4) |
| E3/E3           | 107 (60.5) | 83 (37.2) |
| E3/E4           | 39 (22) | 12 (5.3) |
| No data         | 9 (5) | 111 (50) |

3.2. Evaluation of the Association of the Susceptibility of AD with the Genotype, Allele and Haplotype Frequencies of the IFITM3 Gene in the Korean Population

To investigate genotype and allele frequencies of the IFITM3 gene, we performed direct sequencing in 233 healthy individuals and 177 AD patients and carried out genotyping and HWE analyses. Detailed information on the genotyping results and HWE values in the Korean population is described in Table 2.

Table 2. Comparison of genotype and allele frequencies between healthy controls and Alzheimer’s disease (AD) patients in a Korean population.

| Variants     | Genotype Frequency, n (%) | p-Value | Allele Frequency, n (%) | p-Value | HWE   |
|--------------|---------------------------|---------|-------------------------|---------|-------|
| c.-223C > G  | Controls                  |         |                         |         |       |
| Rs1297064916 | 232 (1) 1 (0) 0 (0)       |         | 465 (1) 1 (0)           |         | 0.9738 |
| Patients     | 177 (1) 0 (0) 0 (0)       |         | 354 (1) 0 (0)           |         |       |
| c.-204G > T  | Controls                  |         |                         |         |       |
| Rs3888188    | 91 (0.39) 109 (0.47) 33 (0.14) |         | 291 (0.62) 175 (0.38)   |         | 0.9686 |
| Patients     | 78 (0.44) 78 (0.44) 21 (0.12) | 0.5546 | 234 (0.66) 120 (0.34)   | 0.2800  | 0.8245 |
| c.-188T > C  | Controls                  |         |                         |         |       |
| Rs6598045    | 146 (0.63) 80 (0.34) 7 (0.03) |         | 372 (0.80) 94 (0.2)     |         | 0.3129 |
| Patients     | 118 (0.67) 53 (0.3) 6 (0.03) | 0.6390 | 289 (0.82) 65 (0.18)    | 0.5161  | 0.9870 |
| c.-181T > C  | Controls                  |         |                         |         |       |
| Rs7478728    | 135 (0.58) 84 (0.36) 14 (0.06) |         | 354 (0.76) 112 (0.24)   |         | 0.8461 |
| Patients     | 118 (0.67) 48 (0.27) 11 (0.06) | 0.1540 | 284 (0.8) 70 (0.2)      | 0.1459  | 0.0532 |
Table 2. Cont.

| Variants | Genotype Frequency, n (%) | p-Value | Allele Frequency, n (%) | p-Value | HWE |
|----------|---------------------------|---------|-------------------------|---------|-----|
|          | AA | AC | CC |          | A | C |       |
| c.-178A > C | Controls 136 (0.58) 83 (0.36) 14 (0.06) | 0.1826 | 355 (0.76) 111 (0.24) | 0.7782 |
| Rs71452596 | Patients 118 (0.67) 48 (0.27) 11 (0.06) | 0.284 (0.8) 70 (0.2) | 0.1665 | 0.0532 |
| c.-175T > C | Controls 136 (0.59) 82 (0.35) 14 (0.06) | 0.2030 | 354 (0.76) 110 (0.24) | 0.1665 | 0.0532 |
| Rs7479267 | Patients 118 (0.67) 48 (0.17) 11 (0.06) | 0.284 (0.8) 70 (0.2) | 0.1785 | 0.0532 |
| c.-128C > T | Controls 229 (0.98) 4 (0.02) 0 (0) | 0.7307 | 350 (0.99) 4 (0.01) | 0.7318 | 0.8792 |
| Rs536077498 | Patients 173 (0.98) 4 (0.02) 0 (0) | - | - | - | - |
| c.-23G > A | Controls 73 (0.3) 116 (0.5) 44 (0.2) | 0.4138 | 262 (0.56) 204 (0.44) | 0.1946 | 0.8228 |
| Rs34481144 | Patients 66 (0.37) 83 (0.47) 26 (0.16) | 0.215 (0.61) 139 (0.39) | 0.1946 | 0.8228 |
| c.-13_-4Del | Controls 232 (1) 1 (1) 0 (0) | 1.0 | 354 (1) 0 (0) | 1.0 | 0.9397 |
| Rs773536963 | Patients 177 (1) 0 (0) 0 (0) | 1.0 | 354 (1) 0 (0) | 1.0 | 0.9397 |

The genotype frequencies of 10 polymorphisms of the IFITM3 gene were in HWE in the healthy controls. To estimate the association between susceptibility to AD and polymorphisms of the IFITM3 gene, we performed case-control association analysis. There were no significant differences ($p > 0.05$) in the genotype or allele distributions of the IFITM3 polymorphisms between healthy individuals and AD patients (Figure 1, Table 2).

Figure 1. Comparisons of the genotype and allele frequencies of rs6598045, rs34481144 and rs12252 single nucleotide polymorphisms (SNPs) of the interferon-induced transmembrane protein 3 (IFITM3)
gene in AD patients and matched healthy controls: (A) Comparison of the genotype frequency of the rs6598045 SNP of the *IFITM3* gene between these two groups; (B) Comparison of the genotype frequency of the rs34481144 SNP of the *IFITM3* gene between these two groups; (C) Comparison of the genotype frequency of the rs12252 SNP of the *IFITM3* gene in these two groups; (D) Comparison of the allele frequency of the rs6598045 SNP of the *IFITM3* gene between these two groups; (E) Comparison of the allele frequency of the rs34481144 SNP of the *IFITM3* gene between these two groups; (F) Comparison of the allele frequency of the *IFITM3* rs12252 SNP between these two groups. N.S.: not significant.

Next, we performed haplotype analysis of 10 polymorphisms within the *IFITM3* gene, namely, c.-223C > G, c.-204G > T, c.-188T > C, c.-181T > C, c.-178A > C, c.-175T > C, c.-128C > T, c.-13.-4DelTTCGCTGGAC, c.42C > T and c.165C > T, in AD patients and healthy controls. All haplotypes showed similar distributions (*p* value > 0.5) in AD patients and healthy controls (Table 3).

| Haplotype          | Frequency (Control) | Frequency (Patients) | *p*-Value |
|--------------------|---------------------|----------------------|-----------|
| CGTTATCwtCC        | 193 (0.545)         | 272 (0.583)          | 0.2705    |
| CTCTCCwTC          | 82 (0.232)          | 90 (0.193)           | 0.1798    |
| CTCTATCwTC         | 49 (0.139)          | 63 (0.135)           | 0.8940    |
| CGCTATCwTC         | 20 (0.056)          | 21 (0.045)           | 0.4569    |
| Others             | 10 (0.028)          | 20 (0.044)           | 0.2677    |

Table 3. Haplotype frequencies of *IFITM3* polymorphisms in the healthy controls and Alzheimer’s disease (AD) patients.

4. Discussion

AD is characterized by the M1 type of disease-associated microglia (DAM) and the elevation of several neuroinflammation-related genes in the brain [23,24]. In a previous study, upregulation of the *IFITM3* protein was also observed with neuroinflammation-related proteins by large-scale proteomic profiling in Aβ-treated microglia cell lines [25]. In addition, a recent study has reported that IFITM3 protein plays a major role in Aβ production via activation of γ-secretase [21]. Thus, the *IFITM3* gene was proposed as a novel therapeutic target for AD. A previous study did not find the association between AD and *IFITM3* SNPs. However, this study was performed with a relatively small sample size and, has not investigated the association of the rs6598045 SNP affecting the expression level of the *IFITM3* gene via TFII-I binding capacity [18,21]. Thus, we performed an association analysis of AD with *IFITM3* polymorphisms in larger populations. However, in the present study, we did not find significant differences in genotype, allele or haplotype frequencies of the *IFITM3* gene between AD patients and matched healthy controls (Tables 2 and 3). In addition, when our data set were stratified by sex and APOE genotype, there were no differences in the genotype, allele or haplotype frequencies of the *IFITM3* gene between AD patients and matched controls (data not shown).

Previous studies have reported that the rs12252 C allele of the *IFITM3* gene was associated with the severity of influenza for the influenza A H1N1 2009 pandemic virus [26–29]. In addition, it was suggested that the severity of influenza A 2009 virus infection varies among ethnic groups based on differences in the distribution of the *IFITM3* rs12252 SNP. The genotype and allele frequencies of rs12252, rs34481144, and rs6598045 SNPs are significantly different between Asian and Caucasian populations. In particular, although the risk of A allele of the rs34481144 SNP exists in less than 1% of East Asian individuals, the A allele of this SNP is present in a high proportion of individuals in European populations and is involved in the severity of pandemic influenza A 2009 virus infection [15,17,18]. Thus, in this study, although the SNPs of the *IFITM3* gene did not show an association with susceptibility to AD, the *IFITM3* SNPs is likely to affect AD phenotypes, including cognitive ability and Aβ accumulation. Thus, further investigation of the effect of the *IFITM3* SNPs on
the phenotype of AD is needed. In addition, since this study investigated only a Korean population, further study is needed to validate the association between \textit{IFITM3} SNPs and susceptibility to AD in other ethnic groups. Furthermore, since the genetic mechanism of the \textit{IFITM3} gene was not related to susceptibility to AD, further investigation of the epigenetic mechanism of the \textit{IFITM3} gene with AD is needed in the future.

5. Conclusions

In this study, we investigated genotype, allele, and haplotype frequencies of the \textit{IFITM3} polymorphisms in AD patients and matched healthy controls. Notably, the genotype, allele, and haplotype distributions of the \textit{IFITM3} polymorphisms did not show significant differences between these two groups. To the best of our knowledge, this is the first report regarding the association analysis of the \textit{IFITM3} gene in AD patients.

Author Contributions: S.-Y.W., Y.-C.K. and B.-H.J. conceived and designed the experiment. S.-Y.W. and Y.-C.K. performed the experiments. S.-Y.W., Y.-C.K. and B.-H.J. analyzed the data. S.-Y.W., Y.-C.K. and B.-H.J. wrote the paper. All authors have read and agreed to the published version of the manuscript.

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Institutional Review Board Statement: A total of blood samples from 177 AD patients were provided by the Pusan National University Hospital Biobank, a member of the Korea Biobank Network. A total of blood samples from 233 healthy Korean subjects were obtained from the Korea Biobank Network at the Centers for Disease Control and Prevention. All experimental procedures were approved according to the guidelines of the institutional review board of Jeonbuk National University and the 1964 Helsinki declaration and its later amendments or comparable ethical standards (approval number: JBNU 2019-03-009).

Informed Consent Statement: Informed consent for all samples was obtained prior to investigation.

Data Availability Statement: All data are available from the corresponding authors upon reasonable request.

Conflicts of Interest: The authors declared no conflict of interest.

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