Genetic Association of Multiple Sclerosis with the Marker rs391745 near the Endogenous Retroviral Locus HERV-Fc1: Analysis of Disease Subtypes

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Abstract

We have previously described the occurrence of multiple sclerosis (MS) to be associated with human endogenous retroviruses, specifically the X-linked viral locus HERV-Fc1. The aim of this study was to investigate a possible association of the HERV-Fc1 locus with subtypes of MS. MS patients are generally subdivided into three categories: Remitting/Relapsing (RRMS), Secondary Progressive (SPMS), and Primary Progressive (PPMS). The association of the HERV-Fc1 locus with the clinical course of MS has not been studied previously. MS patients are clinically subdivided into three categories: Remitting/Relapsing MS (RRMS), Secondary Progressive MS (SPMS), and Primary Progressive MS (PPMS). RRMS and SPMS are generally considered time-wise progressions of the same disease, altogether considered time-wise progressions of the same disease, altogether

Introduction

Multiple sclerosis (MS) is a chronic inflammatory, demyelinating disease of the central nervous system probably caused by interaction of multiple genes and environmental factors [1]. HLA genes are the most strongly associated genes in MS [2], and low Vitamin D level in serum, Epstein Bar Virus infection and smoking are currently the best documented environmental risk factors [3,4].

An involvement of retroviruses has also long been suspected [5]. Originally, this was based on animal models, where other demyelinating diseases could be induced by retroviruses [6,7,8]. However, these animal diseases are all contagious and horizontally transmitted, and MS does not seem to be transferred in this manner. Therefore, some scientists have focused their research on endogenous retroviruses that are vertically transmitted from parents to offspring as part of the chromosomes. Attempts at isolating such viruses from patients with MS have been partial successful [9,10]. The main difficulty of the latter studies, apart from the technical challenges, is the inadequacy of Koch’s postulates for dealing with ubiquitous agents. Koch’s postulates are the criteria traditionally required to be fulfilled in order to establish a causal relationship between a microbe and a disease, and contain the implicit assumption that most people are free of the microbe. The postulates fail, when the microbe is universally present, which is the case with most endogenous viral loci.

Recently, we described the occurrence of MS to be associated with human endogenous retroviruses [11]. The interpretation of these findings did not involve Koch’s postulates, but depended on the theories of genetics. We found association of MS with alleles of the host gene TRIM5 that restricts the replication of a broad variety of retroviruses. This suggested retroviral replication to be involved in this disease, but did not indicate which virus. Hence, we also looked for association of specific endogenous retroviruses with MS. We found an association of MS with markers in and near the endogenous retroviral locus HERV-Fc1, located on the X-chromosome [12]. The associations could be reproduced in several cohorts, encompassing a total of 1060 cases and 2080 controls [11]. The results suggested the involvement of HERV-Fc1 in MS. In addition, we have recently found that expression of HERV-Fc1 RNA is approximately 4-fold higher in plasma from MS patients with recent attacks, compared to plasma from patients in stable remission and from controls [13]. This provides a possible new biomarker and may give clues to further studies of the pathogenic process in MS.

The association of the HERV-Fc1 locus with the clinical course of MS has not been studied previously. MS patients are clinically subdivided into three categories: Remitting/Relapsing MS (RRMS), Secondary Progressive MS (SPMS), and Primary Progressive MS (PPMS). RRMS and SPMS are generally considered time-wise progressions of the same disease, altogether...
called Bout Onset MS (BOMS) representing the vast majority of the patients (85–90%), while PPMS is characterized by a separate disease entity with disease progression from onset (10–15%). In this paper, encompassing 1181 MS cases and 1886 controls from one new (Norwegian) and one previously examined cohort (Danish), we report that BOMS is associated with a marker near the HERV-Fc1 locus, while PPMS disease is not.

Materials and Methods

Ethical approval
Collection of blood samples from MS patients and blood donors for genetic investigations was approved by the Regional Ethics committee (Norway) and local Science Ethical Committees (Denmark). The patients gave written informed consent.

Clinical data
The sub-typing of the patients was based on clinical data using the criteria of Poser [14].

Analysis
DNAs were extracted from the blood by conventional means. The DNAs were genotyped for the rs391745 single nucleotide polymorphism (SNP) located near the X-linked viral locus HERV-Fc1 on a mass spectrometry Sequenom facility using Goldplex reactions (Sequenom, San Diego, CA). The analysis was performed as previously described [11], using the following sequenom primers; rs391745_PCR1: ACGTTGGATGTATAGTATGGCCACCCTC; rs391745_PCR2: ACGTTGGATGGATTCTCAGCATGGACCATC and the rs391745 probe ggtgCAGCATGGACCATCTCTTCAG.

Statistics
The data were analyzed in IBM SPSS version 19 and Excel 2007 (Microsoft, Redmond, WA). For a minority of persons, no subtype or gender information was available, or the typing of rs391745 failed. The main statistic tests performed were \( \chi^2 \) tests. To combine p-values we used Fisher’s method: \( \chi^2 (P, 2n) = -2 \sum \ln (p_i) \), where \( P \) is the combined two-sided p-value and \( p_i \) are the n one-sided p-values to be combined.

Results and Discussion
We analyzed two cohorts of MS patients: 544 cases and 1142 controls from Denmark, and 637 cases and 744 controls from Norway. The Danish cohort was also included in a previous study [11]. Information about individual subtypes of MS had been collected for most patients in both cohorts.

In the previous analysis, the best marker in or near the HERV-Fc1 locus for association with MS was rs391745. Table 1 shows the association of MS and subtypes of MS with rs391745 in the two cohorts. Both cohorts showed an association of the C-allele of the rs391745 SNP in the total MS group (\( p = 0.003 \)). The overrepresentation of C-alleles among cases corresponds to what was observed previously and extends previous findings to a larger set of patients. BOMS also showed a significant association to the C-allele of this SNP (\( p = 0.003 \)). In contrast, PPMS was not associated with any allele of rs391745. This lack of association was present in both cohorts and was emphasized when the cohorts were combined.

To make sure gender issues did not perturb the results, we also performed a gender-stratified analysis of the combined cohorts. The conclusions were the same: BOMS was associated with HERV-Fc1, PPMS was not (Table S1). The p-values were higher, but this was to be expected as the stratified analysis does not contain any cross comparisons: male cases versus female controls and vice versa and thus has less strength.

Three arguments strengthen the tie between MS and HERV-Fc1. First, the association of MS with markers on the X-chromosomal region is limited to an approximately 20 kb region, which includes HERV-Fc1. There are no other known genes in this region [11]. Secondly, the restriction gene TRIM5, which is known to limit retroviral replication, influences the risk of getting MS [11]. Finally, we have found that expression of HERV-Fc1 RNA is four-fold increased in the plasma of MS patients with a recent history of an attack, relative to patients in stable remission and to controls [13]. We take all these data as indications that

| Table 1. Association of rs391745 located near the HERV-Fc1 locus and Multiple Sclerosis. |
|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|
| Cohort          | Status          | C-allele carriers n = 467 | C-allele noncarriers n = 2531 | Frequency of C-allelecarriers | OR\(^2\) (CI95%)\(^2\) (2-sided) | P-value (2-sided) |
| Danish          | All cases       | 102                      | 431                      | 0.19                      | 1.35 (1.03–1.75)               | 0.03             |
|                 | PPMS            | 4                       | 29                       | 0.12                      | 0.79 (0.27–2.26)               | 0.7              |
|                 | BOMS            | 86                      | 359                      | 0.19                      | 1.36 (1.02–1.81)               | 0.03             |
|                 | Controls        | 166                     | 945                      | 0.15                      |                                |                  |
| Norwegian       | All cases       | 105                     | 522                      | 0.17                      | 1.35 (1.00–1.83)               | 0.05             |
|                 | PPMS            | 16                      | 91                       | 0.15                      | 1.18 (0.67–2.10)               | 0.6              |
|                 | BOMS            | 85                      | 411                      | 0.17                      | 1.39 (1.01–1.92)               | 0.04             |
|                 | Controls        | 94                      | 633                      | 0.13                      |                                |                  |
| Combined        | All cases       | 207                     | 953                      | 0.18                      | 1.32 (1.08–1.61)               | 0.003            |
|                 | PPMS            | 20                      | 120                      | 0.14                      | 1.01 (0.62–1.6)                | 0.96             |
|                 | BOMS            | 171                     | 770                      | 0.18                      | 1.35 (1.09–1.67)               | 0.003            |
|                 | Controls        | 260                     | 1578                     | 0.14                      |                                |                  |

Of the persons 70 failed determination of rs391745, and 59 lacked information of MS subtype.
\(^1\)OR: oddsratio for C-allele carriers vs C-allele noncarriers.
\(^2\)CI95%: 95 percent confidence interval.
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development of BOMS may be influenced by the endogenous retrovirus locus HERV-Fc1.

Another corroboration to this conclusion stems from the observation that PPMS has a gender-ratio close to one [15]. This difference in gender ratio was to be expected if HERV-Fc1, located on the X-chromosome, plays a role in BOMS but not in PPMS.

It seems reasonable to speculate on the importance of HERV-Fc1 for disease. If we take the epidemiological approach and assume that rs391745 is the causative variation, we arrive at an etiological fraction of 0.05 to 0.1, i.e., rs391745 is a minor risk determinant for BOMS. Alternatively, if we take the genetic view that rs391745 is a marker in linkage disequilibrium with the true causative variation, the etiological fraction of the latter variation should be higher and the virus could in principle be a primary cause of BOMS. With the latter optimistic view in mind, we are searching for markers with better association to BOMS in and around HERV-Fc1.

Recently, a large genome-wide association study (GWAS) of MS identified an excess of fifty different genes statistically associated to MS [16]. An overrepresentation was observed of genes related to the immune system. This GWAS did not identify association to the HERV-Fc1 region, but this is not in conflict with the HERV-Fc1 data presented here.

In conclusion, this study suggests a molecular distinction of different disease courses in MS by involvement of HERV-Fc1. This warrants further investigations.

**Supporting Information**

**Table S1** Gender-specific p-values and odds-ratios for rs391745 in relation to subtypes of MS.

(DOC)

**Author Contributions**

Conceived and designed the experiments: BAN ABO HFH TP. Performed the experiments: BH KKN MJL. Analyzed the data: BAN. Contributed reagents/materials/analysis tools: ABO HBS HFH EGC. Wrote the paper: BAN BH ABO EGC HFH KKN.

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