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Progressive multifocal leukoencephalopathy in an immunocompetent patient

Nicolien M. van der Kolk¹,a, Peer Arts²,a, Ingeborg W. M. van Uden¹,a, Alexander Hoischen², Frank L. van de Veerdonk³, Mihai G. Netea³ & Brigit A. de Jong¹,4

1Department of Neurology, Radboud University Medical Center, Nijmegen, The Netherlands
2Department of Genetics, Radboud University Medical Center, Nijmegen, The Netherlands
3Department of Internal Medicine, Radboud University Medical Center, Nijmegen, The Netherlands
4Department of Neurology, VU University Medical Center, Amsterdam, The Netherlands

Abstract
Progressive multifocal leukoencephalopathy (PML), a demyelinating disease of the brain, is typically diagnosed in immunocompromised persons. Here, we describe the diagnostic challenge of PML in an apparently immunocompetent patient. Thorough analyses, including cytokine release assays and whole exome sequencing, revealed a deficit in the antiviral interferon gamma production capacity of this patient and compound heterozygous mutations in BCL-2-associated athanogene 3. Interestingly, both factors are associated with reduced expression of John Cunningham virus T-antigen, a protein that plays a key role in viral replication in infected cells. After validation in other patients, our findings may contribute to novel insights into the etiology and possibly treatment of PML.

Introduction
Progressive multifocal leukoencephalopathy (PML) is a destructive demyelinating disease of the central nervous system caused by the John Cunningham virus (JCV), which belongs to the family of polyoma viruses. The onset of the disease is subacute, with a broad range of clinical features. Its course is progressive and often fatal.¹ Carriership of JCV is common among healthy individuals and it remains latent in the kidney, lymphoreticular, or brain tissue. Approximately 70% of adults are seropositive for JCV. Reactivation of the virus, resulting in PML, occurs typically under immunosuppressive conditions.¹ The growing number of immunosuppressive and immunomodulatory therapeutics has resulted into an increased number of individuals at risk for PML.² Although the common denominator in all these conditions is suppression of cellular immunity (either iatrogenic or endogenous), PML may also occur in patients with minimal or occult immune suppression (e.g., idiopathic CD4+ lymphocytopenia, chronic kidney, or liver disease)³ and even occasionally in apparently immunocompetent patients.⁴ Here, we present a case of PML in an apparently immunocompetent patient in whom a
deficit of the interferon γ (IFNγ) pathway, and compound heterozygosity for mutations in BCL2-associated athanogene 3 (BAG3) were identified.

Patient and Methods

A 49-year-old Caucasian male presented with rapidly progressive symptoms of aphasia, dyscalculia, hyperesthesia of the right arm, and headache. Two months after the first symptoms, his clinical condition worsened acutely with an inability to speak and to carry out activities of daily living, apraxia, confusion, and severe headache. His vital signs were normal and neurologic examination confirmed a nonfluent aphasia, dysgraphia, mild facial paresis on the right, a right-sided hemianopsia, hemi-hypaesthesia, and hyperreflexia without extensor plantar responses. He used no medication and had no significant medical history. The family history was unremarkable for neurologic conditions. Laboratory tests showed normal leukocyte counts, including CD3, CD4, and CD8 counts (ratio T4/T8), low infectious parameters, negative serology for human immunodeficiency virus (HIV), Lues, Borrelia, and Herpes Simplex Virus (HSV). Cerebrospinal fluid (CSF) analysis showed normal cell counts, protein and glucose levels, no oligoclonal bands or markers of neurodegeneration (except for a slightly elevated Tau [431 ng/L], normal <300 ng/L), and cytological analysis showed no signs of malignancy. Cerebral MRI showed confluent subcortical white matter T2 hyperintensities predominantly in the left hemisphere, which extended on consecutive MRIs without contrast enhancement or mass effect (Fig. 1). PCR for JCV in serum and CSF was negative.

A stereotactic biopsy of the left frontal lobe showed perivascular lymphocytic infiltration with sporadic enlarged nuclei, which resembled astrocytes. No pathologic oligodendrocyte nuclei were found and immunohistochemical staining for SV-40 (polyomavirus) was negative. However, a positive PCR alone is no confirmation of active virus replication. The PCR for JCV on the biopsy material, however, was positive. The normal immune status and the lack of evidence for an active JCV infection led to the primary diagnosis of tumefactive multiple sclerosis, and the patient was treated consecutively with methylprednisolone intravenously, glatiramer acetate (20 mg/mL once daily), acetaminophen (1000 mg four times daily).
times a day), and plasmapheresis. As deterioration continued the biopsy material was re-examined 7 months after symptom onset, in a tertiary center with extensive expertise on PML. The presence of foam cells, some bizarre astrocytes, and sporadic oligodendrocytes with ground glass appearance combined with the clinical course led to a revision of the diagnosis to PML, and the patient was started on mirtazapine (15 mg once daily). Unfortunately his condition was progressive and he died 9 months after the initial presentation due to cardiopulmonary complications of a bilateral pneumonia. Autopsy confirmed the diagnosis of PML. Macroscopy showed extensive white matter hyperintensities in both hemispheres, with focal gray glass lesions, suggestive for PML. Microscopy showed large white matter hyperintensities with prominent demyelination, and a granular tissue loss. Extensive reactive astrocytosis, and astrocytes with enlarged polymorph hyperchromatic nuclei were found, indicating a viral cytopathogenic effect. A few oligodendrocytes with enlarged nuclei, perivascular foamy macrophages, and some lymphocytic infiltration were seen.

**Immunological assessment**

After the diagnosis of PML an assessment of the capacity of cells isolated from the patient to respond to microbial stimuli was initiated. Peripheral blood mononuclear cells were isolated from blood collected from the patient, and stimulated with the TLR4-ligand lipopolysaccharide (Escherichia coli LPS 10 ng/mL), the TLR3-ligand PolyI:C (5 μg/mL), and a fungal stimulus (heat-killed Candida albicans 10⁵ microorganisms/mL). IFNγ production capacity was measured using an enzyme-linked immunosorbent assay.

**Genetic analysis**

Whole exome sequencing was performed as described earlier. In brief, DNA was isolated from whole blood, enriched with SureSelect v2 exome (Agilent Technologies, Santa Clara, CA) (50 Mb) and sequenced on SOLiD 4 (Life Technologies, Foster City, CA). Variants were called using high stringency settings and annotated with an in-house pipeline containing information from dbSNP134. Variant filtering was applied as previously reported; in brief, we only selected variants affecting coding exons, microRNAs, and canonical splice sites. Subsequently, synonymous variants were filtered out, and only rare variants (frequency of <0.25% in both dbSNP134 and our in-house database containing >2000 exomes) with high quality were reported (Table 1). All genes with rare nonsynonymous variants were systematically checked for involvement role in immunity; based on Gene Ontology terms, mouse knockout phenotypes, information from the Kyoto Encyclopedia of Genes and Genomes or direct interaction with JCV according to NCBI (the latter was performed by searching for JCV in NCBI, and selecting for genes in humans).
<table>
<thead>
<tr>
<th>Selection criteria</th>
<th>Gene name</th>
<th>Gene component</th>
<th>mRNA change</th>
<th>Amino acid change</th>
<th>ExAC allele frequency</th>
<th>ExAC # of alleles/total</th>
<th>SIFT prediction</th>
<th>Polyphen prediction</th>
<th>PhyloP* Grantham score</th>
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<tbody>
<tr>
<td>A(2), B</td>
<td>BAG3</td>
<td>Exon</td>
<td>230C&gt;T</td>
<td>p.P77L</td>
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<td>Benign</td>
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<td>A(2), B</td>
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<td>Exon</td>
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<td>685A&gt;G</td>
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<td>0</td>
<td>Deleterious</td>
<td>Probably damaging</td>
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<td>808T&gt;C</td>
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<td>p.E56X</td>
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<td>0</td>
<td>Deleterious</td>
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<tr>
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<td>Exon</td>
<td>1885C&gt;T</td>
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<td>3/120,732</td>
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<td>Possibly damaging</td>
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<tr>
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<td>2804T&gt;C</td>
<td>p.N9355</td>
<td>0</td>
<td>0</td>
<td>Not scored</td>
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<td>757G&gt;A</td>
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<td>Benign</td>
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<td>494G&gt;A</td>
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<td>N/A</td>
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</table>

Results

Immunological assessment

While cytokine production upon stimulation with LPS and C. albicans was normal compared to eight healthy individuals. IFNγ production induced by PolyI:C was severely impaired in the patient versus control stimulation (Fig. 2). We were unable to perform extensive immunophenotyping of T-, B-, and NK-cell subpopulations as our patient deceased shortly after the diagnosis of PML.

Genetic analysis

Whole exome sequencing provided 5.5 Gb of mapped sequencing data, resulting in an average coverage of the exome of 81.7-fold. Standard variant filtering resulted in 106 rare, nonsynonymous, and canonical splice site variants. Only 17 genetic variants remained after filtering for genes with a possible role in immunity (Table 2). A genome-wide search in NCBI for genes associated with JCV interaction resulted in 21 genes. An overlap between these 21 genes, and the 106 rare variants resulted in one gene for which JCV interaction was described earlier; we found two very rare heterozygous variants in exon 2 of the BAG3. Cosegregation analysis within the family shows that both unaffected parents carry one of the variants in a heterozygous state, and both variants are absent in the healthy brother (Fig. 3). The variants (p. P77L; p. I94F) have been reported at population allele frequencies of 0.029% and 0.076%, respectively. In addition, rare (<1%) homozygous protein-altering variants are only reported in 16 of >60,000 controls.

Discussion

Although rare, similar cases of PML in apparently immunocompetent individuals have been reported. Due to its variable demographics, presenting symptoms, and prognosis, the diagnosis of PML remains a challenge, especially in apparently immunocompetent individuals. The diagnosis can be strengthened by a positive PCR for JCV in the CSF, which has a high sensitivity and specificity. However, the severity of the immunosuppression seems to determine the accuracy of the available tests, as is illustrated by the low JCV DNA copy numbers in the CSF of highly active anti-retroviral therapy treated (HAART) HIV patients, as well as in nearly half of the multiple sclerosis patients treated with the monoclonal antibody natalizumab who were diagnosed with PML. Moreover, JCV PCR in CSF of PML cases with occult, minimal, or no detected immune suppression was often found negative. Also, substantial variability exists with

Figure 3. Family pedigree of our progressive multifocal leukoencephalopathy (PML) case; the unaffected parents were both carrier of one rare BCL2-associated athanogene 3 (BAG3) variant (paternal variant p. P77L, c.230C>T; maternal variants p. I94F, c.280A>T). The compound heterozygosity for these BAG3 variants most likely affects the resistance against John Cunningham virus (JCV) in the index patient. Neither of the two variants is present in the unaffected brother.
might contribute to novel treatment options. IFN-
Interestingly, IFN- was previously reported to inhibit expres-
sion of JCV T-antigen, the major viral regulatory protein.18
In addition, this study showed a significant decrease of JCV
dNA copies in cells upon IFN- treatment in vitro. The
observed IFN- deficit is most likely an important factor in
the pathogenesis of PML in our patient; whole exome
sequencing was performed in order to identify any genetic
defects that could contribute to this deficit. Compound
heterozygous BAG3 variants in our patient are worthy can-
didates for increased susceptibility to PML as BAG3 is impli-
cated in autophagy and apoptosis through intracellular protein control. It was previously shown that overexpression of BAG3 results in a decrease of the JCV replication, and reduced T-antigen expression through autophagic degrada-
tion, thereby controlling the JCV lytic cycle and its interac-
tion with host cells.9 Although it remains speculative, the
observed genetic variants in BAG3 might compromise the
response against JCV T-antigen in our patient, leading to
insufficient IFN- production, and subsequently PML.

Obviously this case report has several limitations and
the results should be interpreted with caution. First,
although the segregation analysis showed an autosomal
recessive inheritance, the family is rather small, which
makes the pathogenicity uncertain. Validating our find-
ings in other cases is therefore crucial, however, due to
the rarity of PML in apparent immunocompetent patients
this remains extremely challenging. We took up this chal-
lenge and performed genetic testing for BAG3 variants in
two previously reported apparent immunocompetent
PML cases.11,12 Unfortunately, no BAG3 variants were
discovered in these patients. Given the number of factors
suggested to be involved in the reactivation of JCV in
order to develop PML and therefore increased hetero-
genosity between these patients, this might not be surpris-
ing.17,19 In addition to the genetic variants in BAG3, we
also provide evidence for low INF- levels in this case.
However, a possible association between the IFN- deficit
and the BAG3 variants remains to be shown.

This case report shows that specific defects in the IFN-
production upon stimulation may be present in PML
patients without known immune deficits. Patients sus-
pected of PML without immunosuppression should, in
addition to regular immunologic screening, be tested for
IFN- deficiency to confirm our findings. In these condi-
tions treatment with IFN- might potentially be an option
for PML, similar to other conditions treated with recombi-
nant IFN-.

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Author Contributions
N. M. K., P. A., and I. W. M. U. drafted the manuscript.
N. M. K., I. W. M. U., F. L. V., and B. A. J. were involved
in the clinical treatment of the patient. P. A. and A. H.
performed the genetic analysis. M. G. N. and F. L. V.
organized the cytokine analysis. M. G. N. and A. H. were
responsible for the laboratory supervision. All authors
reviewed the manuscript.

Conflict of Interest
None declared.

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