

Differential Binding of Autoantibodies to MOG Isoforms in Inflammatory Demyelinating Diseases

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Abstract

Objective

To analyze serum immunoglobulin G (IgG) antibodies to major isoforms of myelin oligodendrocyte glycoprotein (MOG-alpha 1-3 and beta 1-3) in patients with inflammatory demyelinating diseases.

Methods

Retrospective case-control study using 378 serum samples from patients with multiple sclerosis (MS), patients with non-MS demyelinating disease, and healthy controls with MOG alpha-1-IgG positive ($n = 202$) or negative serostatus ($n = 176$). Samples were analyzed for their reactivity to human, mouse, and rat MOG isoforms with and without mutations in the extracellular MOG Ig domain (MOG-ecIgD), soluble MOG-ecIgD, and myelin from multiple species using live cell-based, tissue immunofluorescence assays and ELISA.

Results

The strongest IgG reactivities were directed against the longest MOG isoforms alpha-1 (the currently used standard test for MOG-IgG) and beta-1, whereas the other isoforms were less frequently recognized. Using principal component analysis, we identified 3 different binding patterns associated with non-MS disease: (1) isolated reactivity to MOG-alpha-1/beta-1 ($n = 73$), (2) binding to MOG-alpha-1/beta-1 and at least one other alpha, but no beta isoform ($n = 64$), and (3) reactivity to all 6 MOG isoforms ($n = 65$). The remaining samples were negative ($n = 176$) for MOG-IgG. These MOG isoform binding patterns were associated with a non-MS demyelinating disease, but there were no differences in clinical phenotypes or disease course. The 3 MOG isoform patterns had distinct immunologic characteristics such as differential binding to soluble MOG-ecIgD, sensitivity to MOG mutations, and binding to human MOG in ELISA.

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Glossary

ADEM = acute disseminated encephalomyelitis; **CBA** = cell-based assay; **FACS** = fluorescence-activated cell sorting; **HCs** = healthy controls; **IF** = immunofluorescence; **MOG** = myelin oligodendrocyte glycoprotein; **MOGAD** = MOG-IgG-associated disorders; **MOG-ecIgD** = extracellular MOG immunoglobulin domain; **MOG-IgG** = serum IgG antibodies against MOG; **MS** = multiple sclerosis; **NMOSD** = neuromyelitis optica spectrum disorder; **ON** = optic neuritis; **PCA** = principal component analysis.

Conclusions

The novel finding of differential MOG isoform binding patterns could inform future studies on the refinement of MOG-IgG assays and the pathophysiologic role of MOG-IgG.

Serum immunoglobulin G (IgG) autoantibodies against myelin oligodendrocyte glycoprotein (MOG-IgG) are associated with a spectrum of neurologic diseases including optic neuritis (ON), acute disseminated encephalomyelitis (ADEM), myelitis, seizures, encephalitis, and with brainstem and/or cerebellar involvement.¹⁻⁹ In addition, MOG-IgG appears to be supportive to discriminate these disorders from multiple sclerosis (MS)^{10,11} as reflected by the first diagnostic recommendations for MOG-IgG-associated disorders (MOGADs).^{1,6,12} Furthermore, MOGADs are not only characterized by clinical but also neuropathologic features.^{13,14}

Although most studies use live cell-based assays (CBAs) with the MOG alpha (α) 1 isoform for the measurement of MOG-IgG, previous results were often discrepant, because of different MOG expression vectors (full-length vs extracellular domain), cell lines, read-out systems (immunofluorescence [IF] vs flow cytometry), and other test variations, which aimed to increase specificity and eliminate nonspecific low-titer positivity.^{1,15-17} Several studies attempted to define the molecular epitopes of MOG-IgG with the help of amino acid substitutions or deletions and discovered distinct binding patterns.¹⁸⁻²¹ The most frequent epitopes were located in the loops between the β sheets of the extracellular Ig domain of human MOG (MOG-ecIgD). These findings were extended by other studies showing that only a subset of human MOG-IgG is also reactive to rodent MOG epitopes^{18,19,22,23} and pathogenic in vitro or in vivo.^{19,23,24}

Earlier studies using MOG-ecIgD as an antigen for immunoassays have indicated a lower sensitivity compared with full-length MOG (with the α 1 isoform as the consensus sequence).^{10,25} Although these results indicated binding differences to different MOG variants, no study so far has analyzed antibody responses to different MOG isoforms. Like most other human myelin genes, the MOG gene undergoes extensive alternative splicing and multiple different MOG isoforms have been described in primates, but not in rodents.²⁶⁻²⁹ Although the extracellular Ig domain is present in all these isoforms, they show profound differences in the composition of the intracellular C terminus resulting in alpha (α) or beta (β) isoforms.

The aim of our study was to analyze the serum IgG antibody response to additional MOG isoforms (α 2, α 3, β 1, β 2, and β 3; figure 1) in MOG α 1-seropositive and MOG α 1-seronegative patients and controls. Furthermore, we analyzed whether the use of additional MOG isoforms (either alone or in combinations) improves the specificity of MOG-IgG CBAs.

Methods

Patients and Controls

We performed a retrospective case-control study including serum samples from 378 patients with inflammatory demyelinating diseases and healthy controls (HCs) with known MOG-IgG serostatus recruited from the participating centers for this study (Innsbruck and Vienna, Austria; Verona, Italy; Barcelona, Spain; Dublin, Ireland; Oxford, UK; Zurich, Switzerland; Munich and Lübeck, Germany; and Lyon, France).^{2,10,16,22} We included samples from children (age <18 years) and adults (\geq 18 years) with an aquaporin-4-seronegative non-MS demyelinating disease typically associated with MOG-IgG (n = 214, thereof 191 seropositive MOGAD), MS (n = 64),¹¹ and HCs (n = 100). We also included MOG-IgG-positive MS (n = 8) and HC (n = 3) samples collected at different centers, and therefore, MOG-IgG seropositivity in MS and HCs is overrepresented compared with previous studies. Samples were sent to Innsbruck and reclassified there as MOG-IgG negative (n = 176; 47%; MOG-IgG(H + L) titer 0 to 1:80; MOG-IgG(Fc) negative) or MOG-IgG positive (n = 202; 53%; MOG-IgG(H + L) titer 1:160 to 1:20,480; MOG-IgG(Fc) positive) using a combination of 2 different live IF MOG α 1 CBAs (CBA-IF) as recently described.¹⁶ In addition, 106 follow-up samples of 35 individuals were analyzed with a median observation period of 4.1 years (range 0.8–12.1 years). Because a detailed clinical and radiologic description was not the primary scope of this study, we summarized clinical data of non-MS patients in predominant clinical phenotypes: opticospinal (isolated optic neuritis, myelitis, brainstem syndrome, or a combination of those, a proportion of these patients fulfilled the current diagnostic criteria for neuromyelitis optica spectrum disorder³⁰), cerebral (ADEM according to current diagnostic criteria,³¹

multiphasic disseminated encephalomyelitis, brainstem encephalitis, or encephalitis), or a combination of both (ADEM followed by optic neuritis and opticospinal with cerebral symptoms) and disease courses (monophasic vs recurrent). Clinical and demographic data according to the MOG-IgG serostatus are shown in table e-1, links.lww.com/NXI/A498.

Neuropathologic investigations were available from 6 MOG-IgG-positive cases: 2 with biopsy and autopsy, 3 with biopsy, and 1 with autopsy. Four of these patients had previously been described as case reports and included in a series of patients to describe the pathology of CNS demyelination accompanied by MOG-IgG.¹³ Serum/CSF samples and biopsy/autopsy specimens were referred to the Division of Neuropathology and Neurochemistry, Department of Neurology, Medical University of Vienna for diagnostic purposes and stored at the institutional biobank.

Standard Protocol Approvals, Registrations, and Patient Consents

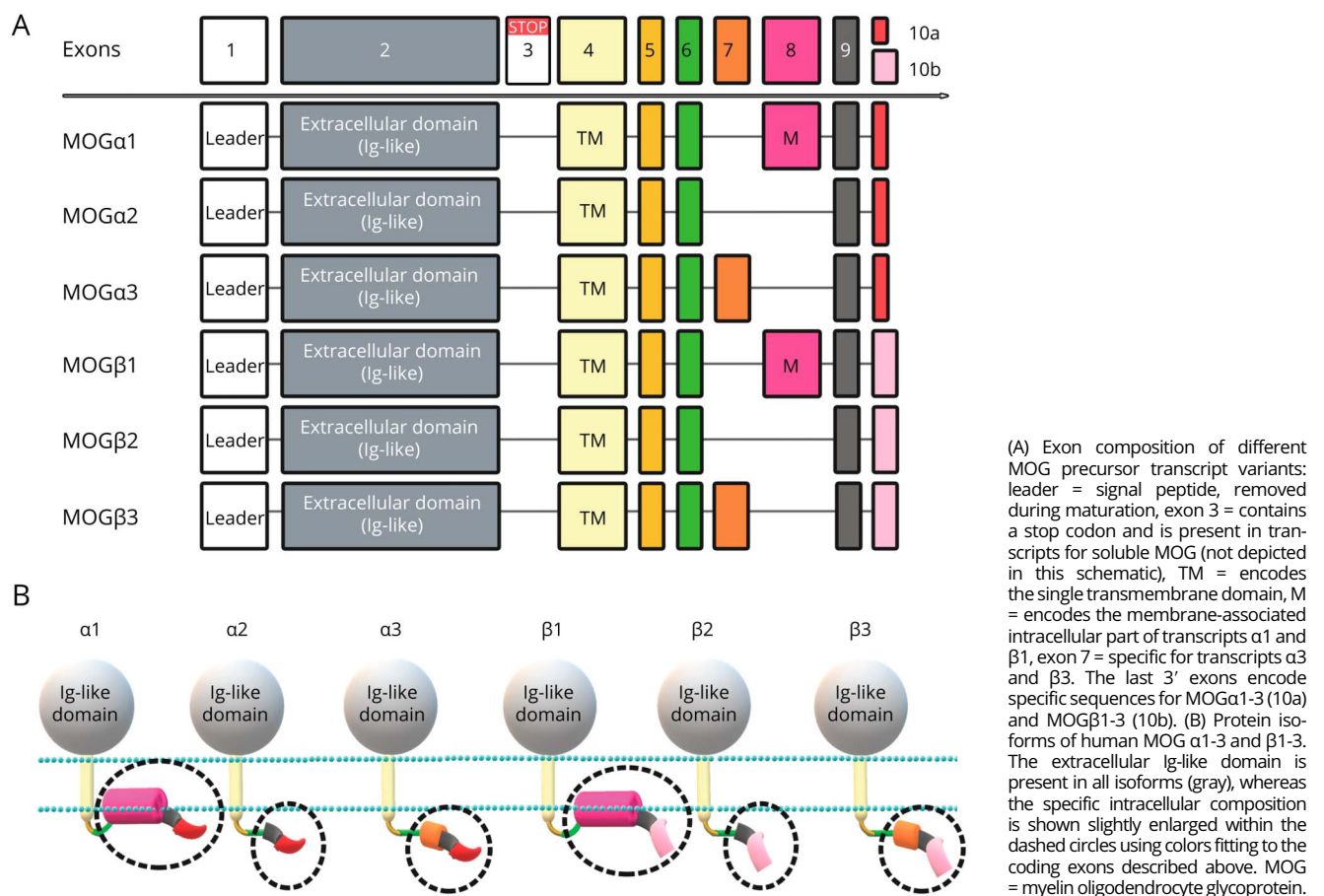
This study was approved by the ethical committees of the Medical University of Innsbruck, Austria (AM3041A and AM4059), Medical University of Vienna (EK 1636/2019 and 1123/2015), University of Oxford, UK (REC 16/SC/0224),

Hospital Clinic of Barcelona, Spain (2010/5680), and University of Zürich, Switzerland (KEK ZH 2013-0001). Twenty serum samples were obtained from the NeuroBioTec—Hospices Civils de Lyon biobank, 10 samples from the Neuropathology-Verona biobank, and 10 samples from the Biobank of LMU Munich, Germany (Biobank 163-16). All patients or their caregivers and controls gave written informed consent. All samples from participating centers were anonymized before sending them to Innsbruck, Austria.

Cloning of MOG Isoform Constructs

The following human MOG isoforms shown in figure 1 were expressed as C-terminal enhanced green fluorescent protein fusion proteins (EGFP) in this study: MOG α 1 (NCBI CCDS34370.1; UniProt Q16653-1), MOG α 2 (NCBI CCDS47394.1; UniProt Q16653-2), MOG α 3 (NCBI CCDS34369.1; UniProt Q16653-3), MOG β 1 (NCBI CCDS4667.1; UniProt Q16653-5), MOG β 2 (NCBI CCDS34367.1; UniProt Q16653-6), and MOG β 3 (NCBI CCDS34366.a; UniProt Q16653-7). The detailed cloning strategies are described in Supplementary methods. In addition, we used pEGFP-N1 (Takara Clontech) expression vectors encoding mouse MOG, rat MOG, and the human MOG α 1 mutants N31Q (N-glycosylation site^{18,20,21,23,32}), P42S (mouse/rat specific, immunodominant human MOG

Figure 1 Splicing Variants and Protein Isoforms of Human MOG α 1-3 and β 1-3



epitope^{18,21,23,32}), E64K (possible binding motif for complement C1q³²), A75S (rat specific^{18,32}), R86Q (mouse/rat specific^{18,32}), and H103A + S104E (epitope of the mouse monoclonal MOG antibody 8-18-C5, important human MOG epitope^{18,21,23,32,33}) as described in Supplementary methods.

Immunoassays for the Detection of MOG-IgG Binding to Different MOG Isoforms

All live CBAs were performed using HEK293 cells transfected with the individual expression vectors encoding the different MOG isoforms described above. Because of the amount of analyses per sample, evaluation of antibody binding to the different isoforms and animal species was performed on the same day to reduce the number of freeze-thawing cycles. All CBAs (live CBA-IF, live CBA-fluorescence-activated cell sorting [FACS], and a commercial fixed CBA-IF from Euroimmun) and ELISAs were performed according to recently published detailed protocols.¹⁶ We used a combination of 2 assays with secondary antibodies to human IgG(H + L) and confirmation by secondary antibodies to human IgG(Fc) to exclude isolated IgM reactivity. Final results were given in titer level (CBA-IF) or binding ratio (CBA-FACS). The binding of serum samples to human, mouse, and rat myelin was analyzed by a tissue-based IF assay as described before.¹⁹

Competitive Binding Assay Using the Soluble Extracellular Domain of MOG

We performed competitive binding assays by CBA-FACS¹⁶ to investigate whether MOG-IgG recognizes soluble human MOG-ecIgD and to determine antibody affinities. Serum samples were diluted according to their FACS median binding ratio to standardize MOG-IgG amounts. Selection of samples was based on the MOG isoform binding pattern observed in live CBA-IF. Diluted serum samples were incubated with increasing amounts (0, 1.6, 3.2, and 9.6 μ M) of soluble human MOG-ecIgD (provided by Euroimmun, Lübeck)¹⁶ for 1 hour at room temperature and spun down at 10,000g for 5 minutes. After this preabsorption step, samples were analyzed by CBA-FACS. Nonlinear regression analysis was used to analyze competitive binding and calculate IC50 values.

Statistical Analysis

The primary hypothesis of this study was that there are differential disease-specific IgG antibody responses to MOG isoforms (α 1, α 2, α 3, β 1, β 2, and β 3). This hypothesis was tested for antibody titers using the Friedman nonparametric test, Spearman's rank nonparametric correlational analyses, and receiver operating curve analysis. Principal component analysis (PCA) was used to classify groups of log2-transformed MOG-IgG titers. This unsupervised, unbiased multivariate analysis approach was used to identify the set of variables (MOG isoforms) accounting for the greatest variation present in the data set. Loading plots were generated to visualize the combination of MOG isoforms responsible for clustering. The analyses of all secondary and other endpoints focused on estimates (common OR; positive and negative predictive values [PPV and NPV]; percentages, medians, and median differences) and 95% CI or

25th to 75th interquartile ranges. Groups were compared using Kruskal-Wallis or χ^2 tests. Statistical significance was defined as 2-sided p value < 0.05 (adjusted for multiple comparisons using Bonferroni's correction). Statistical analyses were performed using IBM SPSS software (IBM SPSS Statistics; Version 26.0. Armonk, NY: IBM Corp.) or GraphPad Prism 9 (GraphPad Software, La Jolla, CA).

Data Availability

The data set used and analyzed during the current study is included in the main text or the supplementary files or is available from the corresponding author on reasonable request. Expression plasmids for the different isoforms will be deposited with Addgene.

Results

Differential Antibody Binding to MOG Isoforms α 1, α 2, α 3, β 1, β 2, and β 3

Because previous studies could not consistently clarify which MOG epitopes are immunodominant, we used a different approach and compared the binding of human MOG-IgG with MOG isoforms α 1, α 2, α 3, β 1, β 2, and β 3. We analyzed serum samples from 378 individuals with MOG-IgG serostatus determined using MOG α 1 live CBA (176 negative and 202 positive) and single transfections of HEK293 cells with the other MOG isoforms. Cell surface expression of isoforms was demonstrated by performing live CBA with serial dilutions of the humanized mouse monoclonal MOG antibody 8-18-C5 showing comparable detection endpoints (figure 2, A and B). Despite small differences in transfection efficiencies, MOG isoforms showed comparable specific surface binding of humanized MOG antibody 8-18-C5 (figures e-2 and e-3, links.lww.com/NXI/A498). Therefore, all extracellular domains of the different MOG isoforms were determined equally available for antibody binding.

Overall, the strongest antibody reactivities were directed against the longest MOG isoforms α 1 (median titer 1:160, 25th–75th percentile 0–640) and β 1 (1:160, 0–640), whereas isoforms α 2 (0, 0–20), α 3 (0, 0–80), β 2 (0, 0–0), and β 3 (0, 0–0) were less frequently recognized ($p < 0.001$). There were strong correlations between α 1 and β 1 (Spearman's ρ 0.990) and between α 2, α 3, β 2, and β 3 (Spearman's ρ 0.677–0.845), whereas the correlations between α 1/ β 1 and α 2/ α 3/ β 2/ β 3 were only moderate (Spearman's ρ 0.427–0.545).

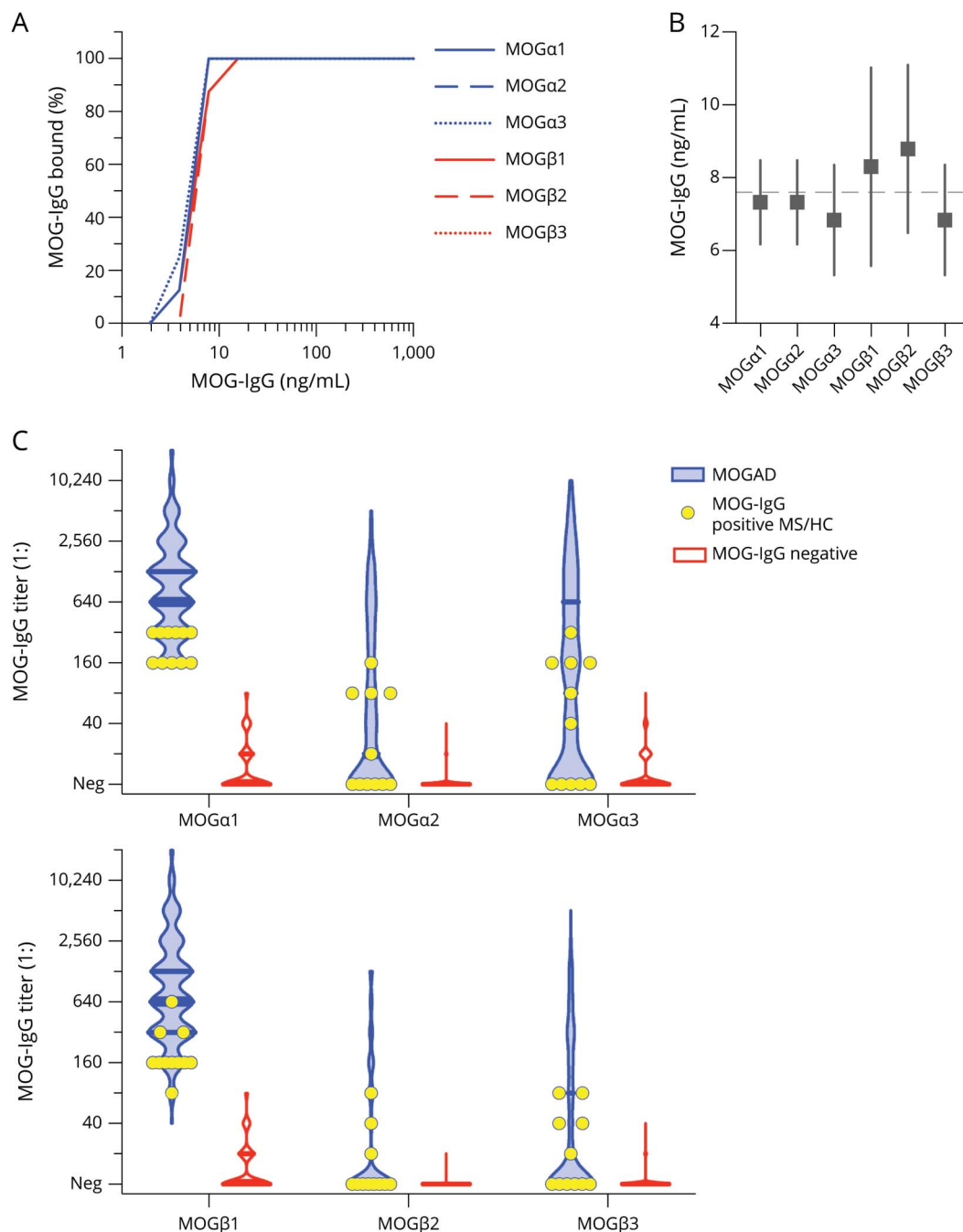
In a next step, we analyzed whether the individual MOG isoforms are useful for discriminating a non-MS demyelinating disease from MS or HC. As can be seen in figure 2C, low-titer antibodies against all MOG isoforms were also present in a few cases with MS and in HCs. The positive and negative predictive values for all MOG isoforms according to titer steps are shown in table e-3, links.lww.com/NXI/A498. The cutoff values associated with a PPV (for MOG-IgG associated with a clinical MOGAD phenotype) of ~95% were \geq 1:160 for MOG α 1 (190/214 non-MS vs 11/164 controls),

$\alpha 3$ (89/214 non-MS vs 4/164 controls), and $\beta 1$ (185/214 non-MS vs 10/164 controls), $\geq 1:40$ for MOG $\alpha 2$ (86/214 non-MS vs 5/164 controls) and $\beta 2$ (52/214 non-MS vs 2/164 controls), and $\geq 1:80$ for MOG $\beta 3$ (58/214 non-MS vs 2/164 controls).

Identification of Distinct Non-MS-Associated MOG Isoform Binding Patterns

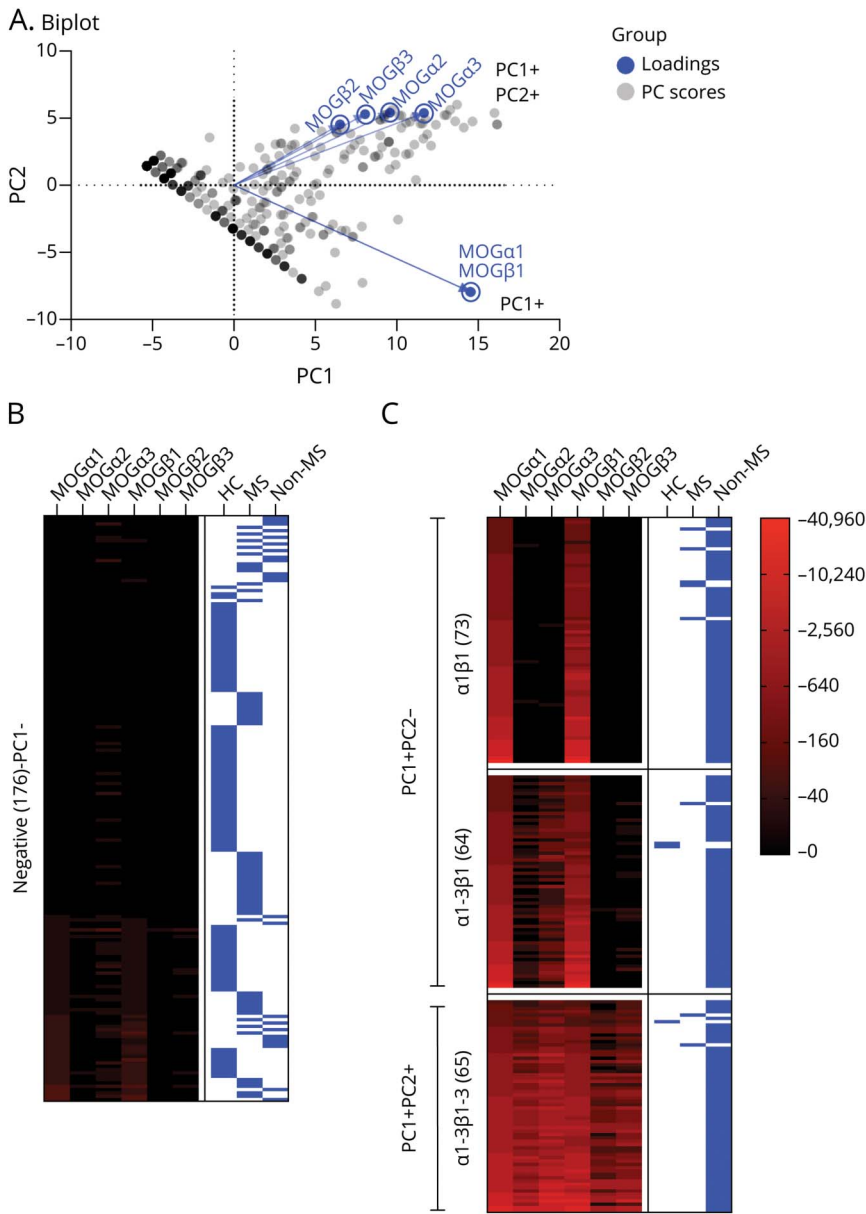
In a next step, we investigated combinations of MOG isoforms using unsupervised PCA to generate a PCA scores plot of all samples and isoforms (figure 3A). The variables driving

Figure 2 Differential Binding of Human MOG-IgG to MOG Isoforms $\alpha 1$, $\alpha 2$, $\alpha 3$, $\beta 1$, $\beta 2$, and $\beta 3$



(A) Serial dilutions of humanized monoclonal mouse MOG antibody 8-18-C5. The graph shows the percentage of 8 replicates determined MOG-IgG positive in live CBA-IF transfected with individual MOG isoforms. (B) Mean sensitivity (with 95%) for MOG-IgG in live CBA-IF using the different MOG isoforms as antigens. The dashed line indicates the overall mean sensitivity. (C) Quantitative binding of MOG-IgG to MOG isoforms $\alpha 1$ -3 (upper panel) and $\beta 1$ -3 (lower panel) for all 378 samples according to their MOG-IgG serostatus (202 positive and 176 negative). Seropositive samples with an aquaporin-4-seronegative non-MS demyelinating disease typically associated with MOG-IgG (MOGAD, $n = 191$) are shown as blue violin plots, and seropositive samples from patients with MS ($n = 8$) or healthy controls ($n = 3$) are shown as yellow circles. Seronegative samples are shown as red violin plots. Medians and interquartile ranges are indicated by the bars within violin plots. CBA = cell-based assay; HCs = healthy controls; IF = immunofluorescence; MOG = myelin oligodendrocyte glycoprotein; MOG-eclgD = extracellular MOG immunoglobulin.

Figure 3 Binding Patterns of Human MOG-IgG to MOG Isoforms $\alpha 1$, $\alpha 2$, $\alpha 3$, $\beta 1$, $\beta 2$, and $\beta 3$



(A) Scatter dot plot showing the principal component (PC) scores of all samples and the loading scores of all MOG isoforms to visualize parameters responsible for clustering. This identified different binding pattern clusters associated with PC1 ($\alpha 1$ and $\beta 1$) and/or PC2 ($\alpha 2$, $\alpha 3$, $\beta 2$, and $\beta 3$). Heatmap of the quantitative results (MOG-IgG titers) for all MOG-IgG-seronegative samples (B) or MOG-IgG-seropositive samples (C) according to their MOG isoform binding pattern (negative, $\alpha 1\beta 1$, $\alpha 1$ - $\beta 1$, and $\alpha 1$ - $\beta 1$ - $\beta 3$). Each column is an individual MOG isoform, and each row is an individual serum sample with MOG isoform binding patterns indicated on the left. MOG-IgG reactivities (titer 1:) are shown in different color intensities (legend with the log scale) and clinical diagnosis of non-MS, MS, and HC according to MOG isoform IgG titers (median with the interquartile range) according to the identified binding patterns. The cutoff value for MOG-IgG positivity ($\alpha 1$ 1:160) is indicated by the dashed blue line. (C) Percentage of patients with clinical diagnosis of non-MS (MOGAD), MS, and HC according to MOG isoform binding patterns. HCs = healthy controls; MOG = myelin oligodendrocyte glycoprotein; MOGAD = MOG-IgG-associated disorders; MOG-IgG = serum IgG antibodies against MOG.

clustering are clearly differentiated in PC1 (all isoforms with strongest contributions of $\alpha 1$ and $\beta 1$; 75.3% of variance, Eigenvalue 28.7) and PC2 ($\alpha 2$, $\alpha 3$, $\beta 2$, and $\beta 3$; 21.2% of variance, Eigenvalue 8.1) together explaining 96.5% of the total variance in the data set. The biplot shown in figure 3A indicates the following major groups: negative for PC1 (seronegative for MOG-IgG), positive for PC1 but negative for PC2, and positive for both PC1 and PC2. This separation into different binding clusters is shown in figure 3 and table 1. The “negative” cluster (PC1 negative, $n = 176$) was associated with MOG-IgG-negative (MOG $\alpha 1$ titer 0-1:80) non-MS, MS, and HC samples. By contrast, the other binding patterns were predominantly associated with non-MS and dominated by either PC1 or PC2: (1) equal binding to $\alpha 1$ and $\beta 1$ and no

binding to $\alpha 2$, $\alpha 3$, $\beta 2$, and $\beta 3$ (pattern $\alpha 1\beta 1$, $n = 73$), (2) equal binding to $\alpha 1$ and $\beta 1$, weaker binding to $\alpha 2$ and $\alpha 3$, and no binding to $\beta 2$ and $\beta 3$ (pattern $\alpha 1$ - $\beta 1$, $n = 64$), and (3) binding to all MOG isoforms (pattern $\alpha 1$ - $\beta 1$ - $\beta 3$, $n = 65$). Three representative CBA-IF stainings for samples assigned to these isoform binding patterns are shown in figure 4.

It is important that these binding patterns were associated with a non-MS demyelinating disease consistent with MOGAD when compared with the MOG-IgG-negative patients and controls (table 1): pattern $\alpha 1\beta 1$ (PPV 94.5%, 95% CI 86.7 to 97.9), pattern $\alpha 1$ - $\beta 1$ (PPV 95.3%, 95.3% CI 87.1 to 98.7), and pattern $\alpha 1$ - $\beta 1$ - $\beta 3$ (PPV 93.8%, 93.9% CI 85.2 to 97.6). The proportion of cases with MOG $\alpha 1$ titers $\geq 1:640$

Table 1 Comparison of Demographic and Clinical Data of All Patients and Controls According to Their MOG Isoform Binding Patterns

	Negative (n = 176)	$\alpha 1\beta 1$ (n = 73)	$\alpha 1-3\beta 1$ (n = 64)	$\alpha 1-3\beta 1-3$ (n = 65)	p Value ^a
MOG$\alpha 1$ (1:)^b	0 (0–20)	640 (320–1,280) ^c	640 (320–1,280) ^c	1,280 (320–2,560) ^c	<0.001
MOG$\alpha 2$ (1:)^b	0 (0–0)	0 (0–0)	20 (0–40) ^{c,d}	640 (160–1,280) ^{c,d,e}	<0.001
MOG$\alpha 3$ (1:)^b	0 (0–0)	0 (0–0)	160 (40–160) ^{c,d}	1,280 (640–2,560) ^{c,d,e}	<0.001
MOG$\beta 1$ (1:)^b	0 (0–20)	640 (320–1,280) ^c	640 (320–1,280) ^c	640 (320–2,560) ^c	<0.001
MOG$\beta 2$ (1:)^b	0 (0–0)	0 (0–0)	0 (0–0)	160 (40–320) ^{c,d,e}	<0.001
MOG$\beta 3$ (1:)^b	0 (0–0)	0 (0–0)	0 (0–20)	320 (80–640) ^{c,d,e}	<0.001
Age (y)^b	36.2 (18.2–45.6)	17.5 (7.2–42.0) ^c	18.1 (6.0–36.9) ^c	23.5 (8.7–43.8)	<0.001
Children (<18 y)^{f,g}	19.9% (14.7–26.4)	52.1% (40.8–63.1) ^c	50.0% (38.1–61.9) ^c	40.0% (29.0–52.1) ^c	<0.001
Females^{f,g}	62.6% (54.5–70.0)	46.6% (35.6–57.9)	50.0% (38.1–61.9)	52.3% (40.4–64.0)	ns
Clinical diagnosis^{f,g}					
Non-MS (MOGAD)	13.1% (8.9–18.8)	94.5% (86.7–97.8) ^c	95.3% (87.1–89.7) ^c	93.8% (85.2–97.6) ^c	<0.001
MS	31.8% (25.4–39.0)	5.5% (2.2–13.2)	1.6% (0.1–8.3)	4.6% (1.3–12.7)	
HC	55.1% (47.7–62.3)	0.0% (0.0–5.0)	3.1% (0.6–10.7)	1.5% (0.1–8.2)	
Clinical phenotype^{f,g}					
Cerebral^h	56.5% (36.8–74.4)	30.4% (20.9–42.1)	31.1% (20.9–43.6)	29.5% (19.6–41.9)	ns
Opticospinalⁱ	43.5% (25.6–63.2)	65.2% (53.4–75.4)	63.9% (51.4–74.8)	62.3% (49.7–73.4)	
Mixed^j	0.0% (0.0–14.3)	4.3% (1.2–12.0)	4.9% (1.3–13.5)	8.2% (3.6–17.8)	
Disease course^{f,g}					
Monophasic	87.0% (67.9–95.5)	55.1% (43.4–66.2)	49.2% (37.1–61.4)	57.4% (44.9–69.0)	ns
Recurrent	13.0% (4.5–32.1)	44.9% (33.8–56.6)	50.8% (38.6–62.9)	42.6% (31.0–55.1)	

Abbreviations: ADEM = acute disseminated encephalomyelitis; MOG = myelin oligodendrocyte glycoprotein; MOGAD = MOG-IgG-associated disorders; NMO = neuromyelitis optica spectrum disorder.

^a p values were adjusted for 12 comparisons using Bonferroni's correction.

^b Median (interquartile range), groups were statistically compared using the Kruskal-Wallis test.

^c Significantly different from the MOG-IgG-negative group.

^d Significantly different from the $\alpha\beta$ group.

^e Significantly different from the $\alpha 1-3\beta 1$ group.

^f Percentage with 95% CI, groups were statistically compared using the χ^2 test.

^g Only patients with an aquaporin-4-seronegative non-MS demyelinating disease typically associated with MOG-IgG (MOGAD).

^h ADEM, MDEM, brainstem encephalitis, or encephalitis.

ⁱ Isolated optic neuritis, myelitis, brainstem syndrome, or a combination thereof (including NMO).

^j ADEM or opticospinal presentation with cerebral involvement.

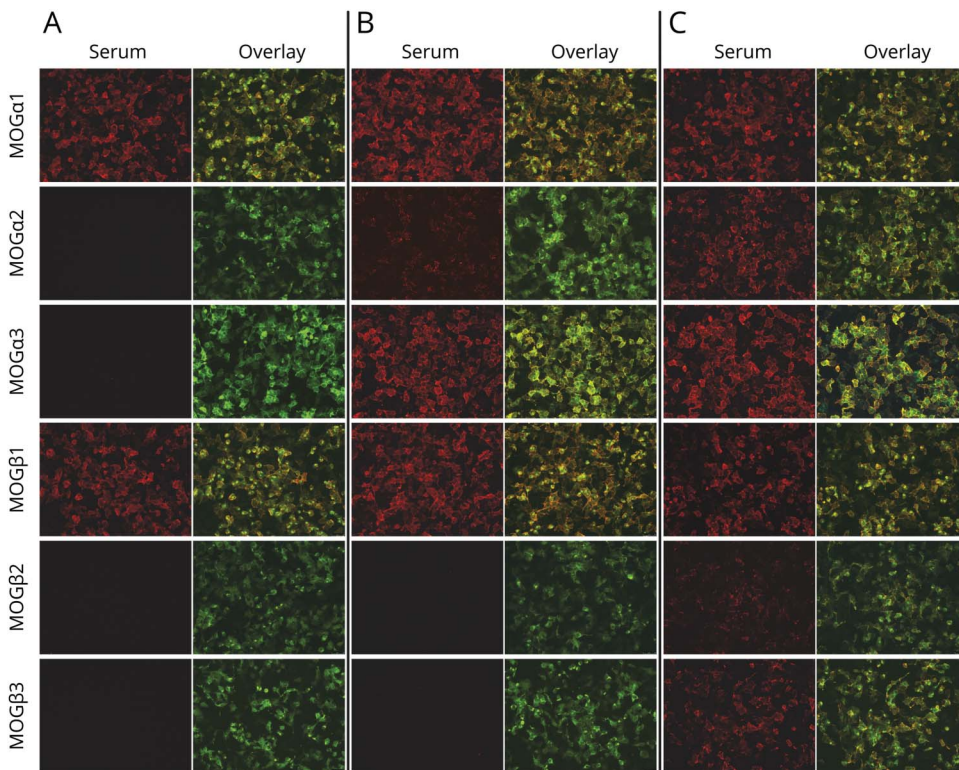
(high-titer MOG-IgG positive) specifically associated with MOGAD¹⁶ was higher in the $\alpha 1-3\beta 1-3$ (74%) than in the $\alpha 1-3\beta 1$ (58%) or $\alpha 1\beta 1$ (59%) binding patterns.

Furthermore, we analyzed 106 available follow-up samples of 35 cases (7 $\alpha 1\beta 1$, 15 $\alpha 1-3\beta 1$, and 13 $\alpha 1-3\beta 1-3$) to assess the temporal stability of these binding patterns. As shown in figure e-4, links.lww.com/NXI/A498, 3 of the 7 (43%) patients with $\alpha 1\beta 1$, 10 of the 15 (71%) patients with $\alpha 1-3\beta 1$, and 9 of the 13 (69%) patients with $\alpha 1-3\beta 1-3$ binding patterns kept their isoform binding pattern. The other patients converted to $\alpha 1\beta 1$ (3) or $\alpha 1-3\beta 1$ (3), or became seronegative (7). There was no conversion from $\alpha 1\beta 1$ to $\alpha 1-3\beta 1-3$ or vice versa.

Associations of Different MOG-IgG Binding Patterns With Non-MS Clinical Phenotype, Disease Course, and Neuropathology

Children with a non-MS disease course were more frequently associated with MOG isoform binding patterns $\alpha 1\beta 1$ and $\alpha 1-3\beta 1$, whereas adults were more frequently associated with $\alpha 1-3\beta 1-3$ or seronegative MOG-IgG (table 1). The percentage of females was comparable between binding patterns. Furthermore, the percentage of patients with different clinical presentations of a non-MS demyelinating disease (cerebral, opticospinal, or mixed presentation) or monophasic or recurrent course at the last follow-up was comparable in all 3 MOG isoform binding patterns. Moreover, there was no association of IgG titers or seropositivity against the individual

Figure 4 Representative MOG Isoform CBA-IF Stainings



Representative live CBA-IF serum stainings for the 3 identified MOG isoform antibody binding patterns (A, $\alpha 1\beta 1$; B, $\alpha 1-3\beta 1$; C, $\alpha 1-3\beta 1-3$). Only specific antibody (red) and overlay images (MOG-transfected cells are shown in green) were used to reduce image size (20 \times magnification). CBA = cell-based assay; IF = immunofluorescence; MOG = myelin oligodendrocyte glycoprotein.

MOG isoforms with demographic or clinical parameters (tables e-4 and e-5, links.lww.com/NXI/A498).

Finally, we analyzed the neuropathologic features at disease/relapse onset and follow-up in the 6 cases with available biopsies and/or autopsies (figure e-5, links.lww.com/NXI/A498). One of these cases (perivenous demyelination) had $\alpha 1\beta 1$ antibodies, 1 had the $\alpha 1-3\beta 1$ binding pattern (perivenous demyelination, conversion to low-titer MOG-IgG at follow-up), and 4 had the $\alpha 1-3\beta 1-3$ binding pattern (1 with perivenous and 3 with confluent demyelination, one of them with intrathecal MOG-IgG).

MOG-IgG Binding Patterns to Different Isoforms Are Associated With a Differential Recognition of MOG Epitopes

We performed a number of additional experiments (reactivity to several mutations in the extracellular Ig domain of MOG $\alpha 1$, competitive binding experiments with soluble MOG-ecIgD, reactivity in CBA-FACS and ELISA, reactivity to fixed MOG expressing cells, and reactivity to human, mouse, and rat MOG in brain tissue) to elucidate the molecular mechanisms underlying the 3 different MOG isoform binding patterns in a subset of samples. Figure 5A shows the differential binding to human MOG $\alpha 1$ mutants, as well as mouse and rat MOG observed in the 3 MOG-IgG binding patterns in relation to MOG $\alpha 1$. The P42S and E64K mutations strongly affected MOG-IgG binding in all 3 binding patterns, with the strongest reduction seen in the $\alpha 1\beta 1$

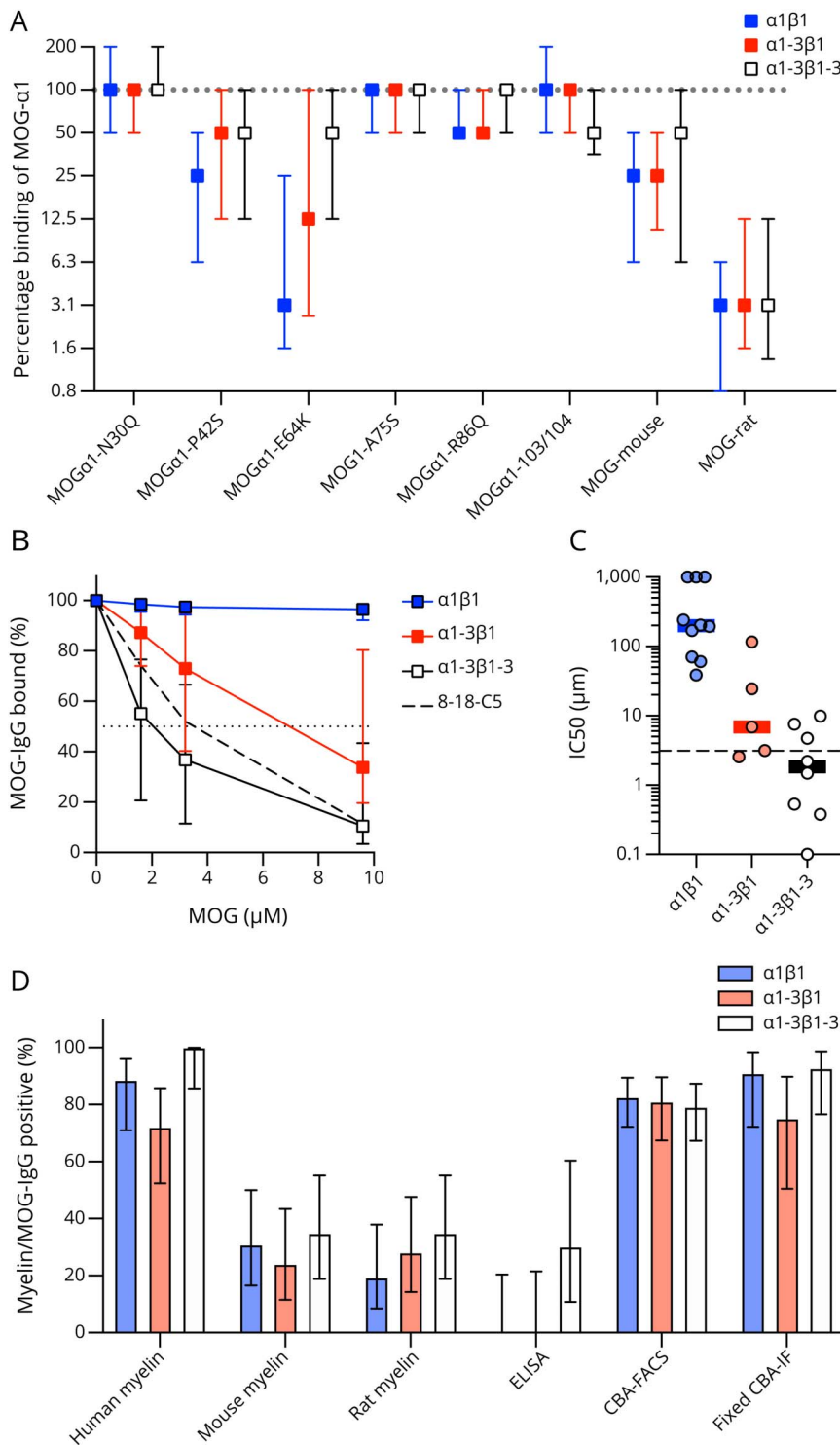
pattern. The N31Q, A75S, R86Q, and H103A + S104E mutations had comparable effects in all binding patterns. Binding to mouse and rat MOG was reduced in all 3 isoform binding patterns.

Next, we analyzed whether the binding to different MOG isoforms can be inhibited by competition with increasing concentrations (0, 1.6, 3.2, and 9.6 μM) of soluble human MOG-ecIgD in a CBA-FACS competitive binding assay. As shown in figure 5B, addition of soluble MOG-ecIgD was able to reduce the binding to MOG $\alpha 1$ in the $\alpha 1-3\beta 1-3$ pattern comparable with the monoclonal MOG antibody 8-18-C5, but not in pattern $\alpha 1\beta 1$. Furthermore, pattern $\alpha 1-3\beta 1-3$ also had similar IC₅₀ and affinities (0–10 μM , figure 5C). By contrast, antibodies of pattern $\alpha 1\beta 1$ did not seem to bind to the soluble MOG-ecIgD. Of interest, pattern $\alpha 1-3\beta 1$ appears to hold a position between the 2. Because median MOG-IgG titers were comparable (1:640 for all 3 binding patterns), we can exclude an antibody concentration effect. Similar findings were obtained using an MOG-IgG ELISA in which $\alpha 1-3\beta 1-3$ antibodies showed better binding than the other patterns (figure 5D). By contrast, we observed no differences for reactivity in CBA-FACS, fixed CBA-IF, and antibody binding to human, mouse, or rat myelin.

Discussion

Our results show that the strongest MOG-IgG responses are directed against the longest MOG isoforms $\alpha 1$ and $\beta 1$, whereas the shorter isoforms $\alpha 2$, $\alpha 3$, $\beta 2$, and $\beta 3$ are less

Figure 5 MOG-IgG Binding Patterns to Different Isoforms are Associated With a Differential Recognition of MOG Epitopes



(A) Binding of MOG-IgG to the human MOGα1 mutants N31Q, P425, E64K, A75S, R86Q, and H103A + S104E (103/104) and to mouse and rat MOG according to their MOG isoform binding patterns (α1β1, n = 64; α1-3β1, n = 43; α1-3β1-3, n = 46). The median differences compared to wild-type MOGα1 are shown as symbols with interquartile ranges (error bars). (B) Competition of binding to MOGα1 in a CBA-FACS assay by soluble MOG-eclgD added in increasing concentrations (0, 1.6, 3.2, and 9.6 μM) for the 3 MOG isoform binding patterns (α1β1, n = 10; α1-3β1, n = 5; α1-3β1-3, n = 8) and for monoclonal antibody 8-18-C5 (dashed line). Squares indicate the median percentage bound, error bars indicate the interquartile ranges, and the value used for the calculation of IC₅₀ (50% binding) is indicated by the dotted line. Groups were statistically compared using repeated measures 2-way analysis of variance. (C) IC₅₀ values for the competition of binding to MOGα1 in a CBA-FACS assay by soluble MOG-eclgD according to MOG isoform binding patterns (α1β1, n = 10; α1-3β1, n = 5; α1-3β1-3, n = 8). Individual data points are shown by scatter dots, and medians and interquartile ranges are indicated by lines and error bars. The IC₅₀ value for monoclonal antibody 8-18-C5 is indicated by the dashed line. (D) Binding of human MOG-IgG according to the 3 MOG isoform binding patterns to human, mouse, and rat myelin in brain sections, as well as in ELISA (human MOG Ig domain), CBA-FACS (human MOGα1), and a commercial fixed CBA-IF (human MOGα1). The percentages of myelin/MOG-IgG-positive samples are shown as bars with 95% CI (error bars). CBA = cell-based assay; FACS = fluorescence-activated cell sorting; IF = immunofluorescence; MOG = myelin oligodendrocyte glycoprotein; MOGα1 = MOG-IgG-associated disorders; MOG-eclgD = extracellular MOG immunoglobulin domain; MOG-IgG = serum IgG antibodies against MOG.

frequently recognized. Using PCA, we could confirm 3 distinct MOG isoform binding patterns specific for patients with MOG-IgG-associated non-MS demyelinating disease (MOGAD): (1) isolated binding to MOG α1 and β1 only (α1β1), (2) a mixed binding pattern with dominant recognition of α1 and β1 and at least 1 additional alpha MOG

isoform (α1-3β1), and (3) binding to all MOG isoforms (α1-3β1-3). Recently, we demonstrated that high-titer positive MOG-IgG (MOGα1 ≥1:640) are consistently positive in all live CBAs and specific for non-MS demyelinating disease, whereas low-titer positive MOG-IgG (MOGα1 1:160–1:320) are more frequently discordant and sometimes also found in

patients with MS and HCs.¹⁶ This observation is also seen in our study with MOG-IgG–positive MS and HC cases found only in the low-titer positive group present in all 3 binding patterns. However, MOG-IgG seropositivity in MS and HCs in our study was overrepresented compared with previous studies.¹

The recognition of different MOG isoforms by MOG-IgG is not associated with relevant differences in clinical presentations, disease course, or neuropathology. Because our study has several limitations such as its retrospective design, the heterogeneous observation period, and missing information on treatment, this needs further investigation in a prospective study. However, demographic and clinical characteristics of our study population are consistent with previous reports.^{1,2,5,6,9,12,22} Our results indicate that $\alpha 1\beta 1$ and $\alpha 1-3\beta 1$ MOG-IgG could be more common in children and $\alpha 1-3\beta 1-3$ MOG-IgG in adults. This observation is consistent with previous findings showing a dominant expression of the isoforms MOG $\alpha 1$ and $\beta 1$ in the human brain, and especially in early development, only these 2 isoforms are detected.^{28,29}

We used a number of additional immunologic investigations to clarify our findings. Only MOG-IgG of the $\alpha 1-3\beta 1-3$, but not the $\alpha 1\beta 1$, binding pattern also bound to soluble MOG-ecIgD with high affinity (20 μM –10 nM), which might explain why immunoprecipitation assays found MOG-IgG less frequently than CBAs^{25,34} and why attempts to isolate human MOG-IgG using affinity binding to MOG-ecIgD were only partly successful.^{21,23} This result is surprising because all MOG-IgG presumably bind to epitopes in the extracellular Ig domain, and therefore, they should compete similarly with MOG-ecIgG. Additional studies are now needed to elucidate the underlying molecular mechanisms of this finding. Furthermore, the fact that only a subset of human MOG-IgG (those of MOG $\alpha 1-3\beta 1-3$ binding pattern) was reactive in MOG ELISA may explain discrepancies found in the literature.^{16,21,25,35} Finally, all 3 MOG isoform binding patterns show different sensitivities to mutations in the extracellular MOG Ig domain: whereas all patterns were sensitive to the P42S mutation associated with an immunodominant MOG epitope^{18,21,23,32} and the E64K mutation in the C'D loop,³⁶⁻³⁸ this effect was even more pronounced in the $\alpha 1\beta 1$ binding pattern. Because the $\alpha 1-3\beta 1$ pattern shares many features with both of the other 2 patterns, it possibly represents a mixture of the other patterns present in polyclonal human serum samples. Further work is now necessary to define the molecular targets of antibodies associated with the identified MOG isoform binding patterns. However, it is a limitation of our study that we were not able to perform these experiments with all samples because of limited availability of samples and reagents.

Our data indicate that these antibody binding patterns are associated with clear structural differences, and therefore, it seems very likely that they represent different antibody clones. The observed MOG isoform binding patterns could

be explained by the differential recognition of MOG dimers or multimers caused by clustering of MOG isoforms.³⁷ This explanation is supported by a recent study demonstrating that the second hydrophobic domain of MOG (only present in $\alpha 1$ and $\beta 1$) enhanced the recognition of the extracellular part of MOG by human MOG-IgG.³⁹ Moreover, MOG-IgG from most patients may require bivalent binding to MOG dimers, whereas a smaller subset of MOG-IgG shows monovalent binding to monomers such as the 8-18-C5 monoclonal antibody.

To conclude, our novel finding of differential MOG isoform binding patterns could explain previous discrepant reports and instruct future studies to improve and refine MOG-IgG antibody assays.

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K. Schanda, P. Peschl, M. Lerch, and B. Seebacher report no disclosures relevant to the manuscript. S. Mindorf has received personal compensation from Euroimmun AG as an employee; N. Ritter has received personal compensation from Euroimmun AG as an employee; M. Probst has received personal compensation from Euroimmun AG as an employee of the Institute for Quality Assurance; H. Hegen has participated in meetings sponsored by, received speaker honoraria or travel funding from Bayer, Biogen, Merck, Novartis, Sanofi-Genzyme, Siemens, and Teva, and received honoraria for acting as a consultant for Biogen and Teva; F. di Pauli has participated in meetings sponsored by, received honoraria (lectures, advisory boards, and consultations) or travel funding from Bayer, Biogen, Celgene, Merck, Novartis, Sanofi-Genzyme, Roche, and Teva. Her institution has received research grants from Roche; E.-M. Wendel has no disclosure to report; C. Lechner served as a consultant for Roche; M. Baumann has no disclosure to report; S. Mariotto received support for attending scientific meetings by Merck and Euroimmun and received speaker honoraria from Biogen; S. Ferrari received support for attending scientific meetings by Shire, Sanofi-Genzyme, Roche, and Euroimmun; A. Saiz reports compensation for consulting services and speaker honoraria from Merck-Serono, Biogen-Idex, Sanofi-Aventis, Teva Pharmaceutical Industries Ltd, Novartis, Roche, and Alexion; M.I.S. Leite reported being

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Continued

Appendix (continued)

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