Prognostic relevance of MOG antibodies in children with an acquired demyelinating syndrome

ABSTRACT

Objective: To assess the prognostic value of MOG antibodies (abs) in the differential diagnosis of acquired demyelinating syndromes (ADS).

Methods: Clinical course, MRI, MOG- and AQP4- abs, and CSF cells and oligoclonal bands (OCB) in children with ADS and 24 months of follow-up were reviewed in this observational prospective multicenter hospital-based study.

Results: Two hundred ten children with ADS were included and diagnosed with acute disseminated encephalomyelitis (ADEM) (n = 60), neuromyelitis optica spectrum disorder (NMOSD) (n = 12), clinically isolated syndrome (CIS) (n = 101), and multiple sclerosis (MS) (n = 37) after the first episode. MOG-abs were predominantly found in ADEM (57%) and less frequently in NMOSD (25%), CIS (25%), or MS (8%). Increased MOG-ab titers were associated with younger age (p = 0.0001), diagnosis of ADEM (p = 0.005), increased CSF cell counts (p = 0.011), and negative OCB (p = 0.012). At 24-month follow-up, 96 children had no further relapses. Thirty-five children developed recurrent non-MS episodes (63% MOG-, 17% AQP4-abs at onset). Seventy-nine children developed MS (4% MOG-abs at onset). Recurrent non-MS episodes were associated with high MOG-ab titers (p = 0.0003) and older age at onset (p = 0.024). MS was predicted by MS-like MRI (p < 0.0001) and OCB (p = 0.007). An MOG-ab cutoff titer ≥1:1,280 predicted a non-MS course with a sensitivity of 47% and a specificity of 100% and a recurrent non-MS course with a sensitivity of 46% and a specificity of 86%.

Conclusions: Our results show that the presence of MOG-abs strongly depends on the age at disease onset and that high MOG-ab titers were associated with a recurrent non-MS disease course.

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GLOSSARY

abs = antibodies; ADEM = acute disseminated encephalomyelitis; ADEMON = acute disseminated encephalomyelitis with optic neuritis; ADS = acquired demyelinating syndromes; CIS = clinically isolated syndrome; EDSS = expanded disability score; FLAIR = fluid-attenuated inversion recovery; LETM = longitudinal extensive transverse myelitis; MDEM = multiphasic acute disseminated encephalomyelitis; MOG = myelin oligodendrocyte glycoprotein; MS = multiple sclerosis; NMOSD = neuromyelitis optica spectrum disorder; OCB = oligoclonal bands; ON = optic neuritis; OND = other neurologic diseases; TM = transverse myelitis.

The differential diagnosis of children with acquired demyelinating syndromes (ADS) encompasses monophasic diseases such as acute disseminated encephalomyelitis (ADEM), clinically isolated syndromes (CIS), or potentially relapsing forms such as multiple sclerosis (MS). Early diagnosis is important because of other disorders mimicking a first episode of ADS and different treatment options. In this context, antibodies (abs) to myelin oligodendrocyte glycoprotein (MOG) have regained attention. Serum MOG-abs can be detected in children with monophasic ADEM with MOG-abs that drop to undetectable levels.\(^1\)\(^-\)\(^3\) MOG-abs have further been detected in aquaporin (AQP)-4- ab-seronegative NMOSD, recurrent optic neuritis (ON), transverse myelitis (TM),\(^4\)\(^-\)\(^7\) and multiphasic acute disseminated encephalomyelitis (MDEM), but less frequently in adults with these disorders.\(^2\)\(^-\)\(^8\)\(^-\)\(^10\) Two recent studies revealed that MOG-abs are nearly exclusively detectable in children with monophasic or relapsing events other than MS.\(^11\)\(^-\)\(^12\)

Author affiliations are provided at the end of the article.

Go to Neurology.org for full disclosures. Funding information and disclosures deemed relevant by the authors, if any, are provided at the end of the article.
In the current study, we investigated the prevalence and predictive value of MOG-abs in children with a first ADS in combination with clinical and MRI features at onset and outcome after 2 years.

METHODS Between 2009 and 2014, serum samples of 302 children with a suspected ADS were sent to us from 60 different medical centers in Germany, Austria, Italy, Switzerland, Turkey, and Canada for testing of MOG- and AQP4- abs and included in an ongoing prospective study. Two hundred ten children with a data set including a diagnosis of ADS, clinical presentation, CSF studies (cell count, oligoclonal bands [OCB]), cerebral MRI from the first episode, MOG- and AQP4- abs status up to 3 months after first episode, treatment history, final diagnosis, and clinical outcome with an expanded disability score (EDSS) after at least 24 months were selected. Clinical data at onset and follow-up were obtained in the majority of cases with a standardized questionnaire or a medical discharge summary from the referring physician.

Ninety-two children were excluded for the following reasons: (1) 2 children with ADEM died at disease onset; (2) 14 children were diagnosed with other neurologic diseases (OND) (e.g., limbic encephalitis); (3) 11 children presented as having ADS but were later diagnosed with OND; (4) in 23 patients, the initial serum sample was not taken during the first 3 months; (5) in 2 children, necessary data from disease onset were not available; (6) 40 children were lost to follow-up (figure 1). Based on the initial clinical information, radiologic evaluation and AQP4- abs status children were assigned to one of the following diagnoses based on the International Paediatric MS Study Group (IPMSG) recommendations5; (1) 60 patients with ADEM; (2) 101 patients with CIS divided further into 6 subcategories (isolated ON, n = 42; bilateral ON, n = 6; transverse myelitis, n = 20; brainstem, n = 7; cerebral, n = 11; polysymptomatic, n = 15); (3) 57 patients were diagnosed with MS because of radiologic evidence of dissemination in space and time; (4) 12 patients with NMOSD (figure 1 and table e-1 at Neurology.org). MRI studies were assessed by at least 2 reviewers unaware of the clinical presentation and MOG- abs status (M.B., T.G., and G.K.) including the following sequences: cranial MRI of all patients in T2-, fluid-attenuated inversion recovery (FLAIR)- axial, T2-sagittal, T1-axial with contrast-medium sequences in addition to spinal T2- and T1-sagittal with contrast-medium sequences available from 160/210 children. Clinical follow-up including EDSS, presence of cognitive dysfunction (e.g., attentional problems), or epilepsy was obtained through medical records or a standardized questionnaire at least after 24 months. Demographic and clinical findings of 63/210 children from this cohort were reported already in 4 studies focusing on the presence of MOG- abs in NMOSD (n = 21 and n = 3016), ADEM (n = 281), and MDEM (n = 3). We decided to include these children because of new clinical data (e.g., further relapses and additional serum samples).

RESULTS Patients. Children with ADS and a complete data set were included (figure 1). The cohort of 210 children consisted of 120 female and 90 male patients with a median age at onset of 12.0 years (range 0.5–17). Sixty patients were diagnosed with ADEM, 101 with CIS, and 12 with NMOSD. Thirty-seven patients with CIS had radiologic evidence of dissemination in space and time and were thus diagnosed with MS (figure 1 and table e-1). After 24 months, the final diagnosis remained ADEM in 46/60 children. Fourteen of 60 children had further events and were diagnosed with NMOSD (n = 1), MS (n = 2), MDEM (n = 8), or acute disseminated encephalomyelitis with optic neuritis (ADEMON) (n = 3). Fifty-four of 101 children with CIS had further demyelinating episodes: 11 children with ON had further episodes of ON and were diagnosed with recurrent ON. Three children with ON or longitudinal extensive transverse myelitis (LETM) had additional episodes of ON or LETM, increasing the number of children with NMOSD to 16 after 2 years. Forty children with CIS were diagnosed with MS due to clinical relapses (n = 35) or new FLAIR/T2 lesions on follow-up MRI (n = 5) (figure 1). Comparison of clinical and immunologic features according to MOG- abs status at baseline. Serum MOG- abs with a titer ≥1:160 were detected in 65/210 (31%) patients with a median titer of 1:1,280 (range 1:160–1:40,960). Patients with high MOG- ab titers had a lower age at onset (Spearman \( r = -0.416, p < 0.0001 \)). Accordingly, MOG- ab titers were highest in
MOG-ab titers were highest in children with ADEM (figure 2). Because previous studies have analyzed smaller patient cohorts using explorative statistical methods, we decided to analyze the association of MOG-ab titer levels (0–80, 160–640, and $1:1,280$) using a multivariate model (table 1). High MOG-ab titers were associated with younger age ($p = 0.00001$), diagnosis of ADEM ($p = 0.005$), absence of OCB ($p = 0.012$), and higher CSF cell counts ($p = 0.011$). By contrast, diagnosis of CIS with MS-like MRI or MS was associated with MOG-ab seronegativity or low titers. MOG-ab–positive children with CIS and MS-atypical MRI exclusively presented with ON ($n = 18$) or TM ($n = 7$) (table 1).

Prognostic parameters for a final diagnosis of MS or other forms of recurrent demyelinating syndromes after 24 months. Next, we studied the prognostic value of baseline markers for final disease classification in the entire MOG-ab–positive and MOG-ab–negative cohort (table 2 and table e-2). After 24 months, 96 children still had a monophasic diseases course (42% MOG-abs at onset), 35 children developed further demyelinating episodes other than MS (63% MOG-abs and 17% AQP4-abs at onset), and 79 children were diagnosed with MS (4% MOG-abs at onset). Multinominal logistic regression analysis indicated that a recurrent non-MS course was predicted by higher MOG-ab titers ($p = 0.0003$) and older age at onset ($p = 0.024$) compared with the monophasic group (table 2). By contrast, MS was predicted by an...
MS-like MRI ($p = 4 \times 10^{-7}$) and OCB at onset ($p = 0.007$) compared with the monophasic group with no significant role of MOG-abs. Of note, 28/35 (80%) patients with a recurrent non-MS course had abs to either MOG (63%) or AQP4 (17%), and 27/35 (77%) patients were females (table 2). Thus, patients with a higher age (median 10 years), female sex, MOG- or AQP4-abs, and MS-atypical MRI at onset have a high risk of developing a non-MS-like recurrent disease course independent from the initial diagnosis. Furthermore, these patients had a higher risk for ON during the course of disease (26/35, 74%) compared with patients with a monophasic disease course (34/96, 35%) or MS (19/79, 24%). After 24 months, no significant difference in the EDSS or other clinical residuals was noted between the different groups (tables e-2 and e-4). Patients in the MOG-ab-negative recurrent non-MS group (most of them with NMOSD with AQP4-abs) and in the group with MOG-ab titers $\geq 1,280$ had the shortest time to relapse. Of 35 (40%) patients in the recurrent non-MS group were treated with IV immunoglobulin ($n = 3$), rituximab ($n = 3$, AQP4-positive NMOSD), or azathioprine ($n = 4$) compared with 3/96 (3%) in the monophasic group or 60/79 (76%) in the MS group.

Finally, we analyzed whether different MOG-ab cutoff titer levels provided prognostic information on the disease course after 2 years. Although our currently used MOG-ab cutoff value of $\geq 1:160$ had the highest sensitivity for predicting a non-MS course or a recurrent non-MS course after 2 years, the corresponding specificities were modest (table 3). By contrast, a cutoff value of $\geq 1:1,280$ had a lower sensitivity for predicting a non-MS course or a recurrent non-MS course, but the corresponding specificities were much better.

**Disease course of MOG-ab-seropositive patients over 24-month follow-up.** Forty of 65 (61%) MOG-ab-seropositive children had a monophasic disease course. Twenty-two children (34%) developed a recurrent non-MS disease course and only 3 children (5%) were diagnosed with MS (table e-3). MOG-ab-positive patients with a recurrent non-MS disease course were diagnosed with MDEM or ADEM-ON ($n = 11$), followed by recurrent ON ($n = 8$) and NMOSD ($n = 3$). Sixteen of 22 (73%) patients with a recurrent disease course had ON at beginning or during follow-up compared with 13/40 (32%) patients with a monophasic disease and 2/3 (66%) with MS. Twenty-one of 22 MOG-ab-positive patients with a recurrent non-MS
entered at the first step (enter method). R2

Groups were statistically compared using ordinal regression analysis with all parameters

verse myelitis.

disorder; OCB

tein; MS

0.289 (McFadden); model

904

h Sixteen missing cases were included in the analysis with values entered as samples means

g Six missing cases were included in the analysis with values entered as samples means.

d Symptoms: brainstem (6), cerebral (11), ON (6), polysymptomatic (12), and TM (2).

c Symptoms: ON (8) and TM (4).

b Diagnosis at onset and MRI at onset were analyzed together because they were highly

correlated, and MRI was used to establish the diagnosis of MS.

Abbreviations: ab = antibody; ADEM = acute disseminated encephalomyelitis; AQP-4 = aquaporin-4; CIS = clinically isolated syndrome; MOG = myelin oligodendrocyte glycoprotein; MS = multiple sclerosis; neg = seronegative; NMOSD = neuromyelitis optica spectrum disorder; OCB = oligoclonal IgG bands; ON = optic neuritis; pos = seropositive; TM = transverse myelitis.

Groups were statistically compared using ordinal regression analysis with all parameters entered at the first step (enter method). R2 = 0.378 (Cox and Snell), 0.468 (Nagelkerke), 0.289 (McFadden); model χ² = 99.73; p = 5 × 10−18.

*Median (range).

Symptoms: brainstem (1), ON (24), polysymptomatic (3), and TM (11).

Symptoms: ON (8) and TM (4).

Symptoms: ON (10) and TM (3).

Symptoms: brainstem (6), cerebral (11), ON (6), polysymptomatic (12), and TM (2).

Six missing cases were included in the analysis with values entered as samples means.

Sixteen missing cases were included in the analysis with values entered as samples means (data shown as median with range).

disease course had a clinical relapse after 6 months (1–13 months) and 2.2 relapses (1–5 relapses) in the first 2 years, which was not different compared with other recurrent groups (table e-3). Of note, an MS-like MRI (e.g., lesions perpendicular to the long axis of the corpus callosum) was absent in the monophasic and recurrent non-MS patients.19 MOG-ab titers at presentation were similar in MOG-ab–positive children with a monophasic vs a recurrent non-MS disease course and lower in patients with MS.

Correlation of MOG-ab follow-up titers with the clinical disease course. Finally, we analyzed 218 follow-up serum samples from 75/145 (52%) MOG-ab–negative patients and 51/65 (78%) MOG-ab–positive patients (cutoff ≥1:160). MOG-ab–negative patients did not develop MOG-ab over time. Overall, 36/51 (71%) MOG-ab–positive patients still had MOG-ab at final follow-up with an overall decline of the median titer from 1:1,280 (range 160–40,960) at onset to 1:160 (range 0–5,120). MOG-ab titers dropped from onset to follow-up in all groups, with the strongest effect seen in the monophasic and MS groups, but remained highest in the recurrent non-MS group (figure 2). When using a cutoff value of ≥1:1,280 as calculated above, 33/126 (26%) patients were MOG-ab positive at onset, but only 7/126 (6%) still had MOG-ab at follow-up. Nineteen of 48 monophasic patients had MOG-ab ≥1:1,280 at onset, but only 1 patient (2%) had persisting abs, whereas 15/26 patients with a recurrent non-MS course had MOG-abs at onset (58%), 6 of them (23%) with persisting abs. None of the 52 patients with MS had MOG-abs ≥1:1,280 at onset or follow-up.

**DISCUSSION** We report the frequency of serum MOG-ab in a cohort of 210 children presenting with ADS prospectively followed for at least 2 years. We confirm previous findings such as the presence of MOG-ab in one-third of all children with ADS and that MOG-ab are associated with a non–MS-like disease course.11,12 As in previous studies, we observed a strong correlation between age and MOG-ab seropositivity at disease onset,5,11–17,18 revealing that, in particular, children younger than 10 years with an acute non–MS-like episode harbor MOG-ab (table e-4). Furthermore, we show that a subgroup of children continued to have further demyelinating episodes such as recurrent ON, MDEM, ADEMON, or NMOSD characterized by the presence of MOG-abs and less often AQP4-abs, higher age at onset, and female predominance. Recurrent episodes in most cases were accompanied by ON, occurred in the context of persisting, high-titer MOG-ab, and in the first year after the initial event. The characteristics of this subgroup of patients are similar to those recently reported in adult MOG-ab–positive patients.19,20 To evaluate what characterizes children with MOG-ab who develop a relapsing non-MS-like disease course, we compared them with children with a monophasic disease course. The initial features of both groups had notable differences: children with monophasic disease course were younger, more often males, and less often had involvement of optic nerves. At initial presentation, antibody titers in children with transient MOG-abs were similar to those with persistent MOG-abs but declined in the following months. Taking all aspects into account, 2 subgroups with MOG-abs emerge, as already suggested in a smaller cohort of pediatric patients17: One subgroup characterized by children with a median age at onset of 4.5 years and equal sex distribution is primarily diagnosed with ADEM and less often with CIS associated with a monophasic course and transient MOG-abs; the

**Table 1** Demographic and clinical parameters at sampling associated with serum MOG-ab titers in 210 pediatric patients with acquired demyelinating diseases

<table>
<thead>
<tr>
<th>No. of cases</th>
<th>MOG-ab neg (0–1:80)</th>
<th>MOG-ab pos (1:160–640)</th>
<th>MOG-ab pos (≥1:1,280)</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Females, n (%)</td>
<td>85 (59)</td>
<td>11 (46)</td>
<td>24 (58)</td>
<td>0.353</td>
</tr>
<tr>
<td>Age, y</td>
<td>13 (1–17)</td>
<td>9 (3–15)</td>
<td>5 (0–17)</td>
<td>0.000001</td>
</tr>
<tr>
<td>Diagnosis, n (%)</td>
<td>39 (27)</td>
<td>12 (50)</td>
<td>13 (32)</td>
<td>0.005</td>
</tr>
<tr>
<td>CIS + atypical MRI</td>
<td>37 (25)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0.999</td>
</tr>
<tr>
<td>MS</td>
<td>34 (23)</td>
<td>3 (12)</td>
<td>0 (0)</td>
<td>0.012</td>
</tr>
<tr>
<td>Polysymptomatic, n (%)</td>
<td>87 (60)</td>
<td>12 (50)</td>
<td>30 (73)</td>
<td>0.136</td>
</tr>
<tr>
<td>Optic neuritis, n (%)</td>
<td>6 (4)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0.999</td>
</tr>
<tr>
<td>AQP4-abs, n (%)</td>
<td>77 (53)</td>
<td>5 (21)</td>
<td>2 (5)</td>
<td>0.012</td>
</tr>
<tr>
<td>CSF cells/µL</td>
<td>8 (0–138)</td>
<td>7 (0–338)</td>
<td>22 (0–338)</td>
<td>0.011</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Value</th>
<th>p Value</th>
</tr>
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</table>
| 1:1,280 at onset, but only 1 patient (2%) had persisting abs, whereas 15/26 patients with a recurrent non-MS course had MOG-abs at onset (58%), 6 of them (23%) with persisting abs. None of the 52 patients with MS had MOG-abs ≥1:1,280 at onset or follow-up.

**DISCUSSION** We report the frequency of serum MOG-ab in a cohort of 210 children presenting with ADS prospectively followed for at least 2 years. We confirm previous findings such as the presence of MOG-ab in one-third of all children with ADS and that MOG-ab are associated with a non–MS-like disease course.11,12 As in previous studies, we observed a strong correlation between age and MOG-ab seropositivity at disease onset,5,11–17,18 revealing that, in particular, children younger than 10 years with an acute non–MS-like episode harbor MOG-ab (table e-4). Furthermore, we show that a subgroup of children continued to have further demyelinating episodes such as recurrent ON, MDEM, ADEMON, or NMOSD characterized by the presence of MOG-ab and less often AQP4-abs, higher age at onset, and female predominance. Recurrent episodes in most cases were accompanied by ON, occurred in the context of persisting, high-titer MOG-ab, and in the first year after the initial event. The characteristics of this subgroup of patients are similar to those recently reported in adult MOG-ab–positive patients.19,20 To evaluate what characterizes children with MOG-ab who develop a relapsing non-MS-like disease course, we compared them with children with a monophasic disease course. The initial features of both groups had notable differences: children with monophasic disease course were younger, more often males, and less often had involvement of optic nerves. At initial presentation, antibody titers in children with transient MOG-abs were similar to those with persistent MOG-abs but declined in the following months. Taking all aspects into account, 2 subgroups with MOG-abs emerge, as already suggested in a smaller cohort of pediatric patients17: One subgroup characterized by children with a median age at onset of 4.5 years and equal sex distribution is primarily diagnosed with ADEM and less often with CIS associated with a monophasic course and transient MOG-ab; the
Table 2  Predictive factors at disease onset for a recurrent non-MS disease course or MS after 24 months in 210 pediatric patients with acquired demyelinating diseases

<table>
<thead>
<tr>
<th></th>
<th>Monophasic</th>
<th>Recurrent non-MS</th>
<th>MS</th>
</tr>
</thead>
<tbody>
<tr>
<td>No. of cases, n (%)</td>
<td>96 (45.7)</td>
<td>35 (16.7)</td>
<td>79 (37.6)</td>
</tr>
<tr>
<td>Females, n (%)</td>
<td>47 (49)</td>
<td>27 (77)</td>
<td>46 (58)</td>
</tr>
<tr>
<td>Age at onset, y*</td>
<td>7 (0.5-17)</td>
<td>10 (2-17)</td>
<td>14 (1-17)</td>
</tr>
<tr>
<td>MOG-abs at onset, n (%)</td>
<td>15 (16)</td>
<td>6 (17)</td>
<td>3 (4)</td>
</tr>
<tr>
<td>AQP4-abs at onset, n (%)</td>
<td>0 (0)</td>
<td>6 (17)</td>
<td>0 (0)</td>
</tr>
<tr>
<td>CSF OCB at onset, n (%)</td>
<td>15 (16)</td>
<td>3 (9)</td>
<td>6 (15)</td>
</tr>
<tr>
<td>MS-typical MRI at onset</td>
<td>4 (4)</td>
<td>0 (0)</td>
<td>71 (90)</td>
</tr>
<tr>
<td>Polysymptomatic onset</td>
<td>62 (65)</td>
<td>20 (57)</td>
<td>47 (60)</td>
</tr>
<tr>
<td>Optic neuritis at onset</td>
<td>34 (35)</td>
<td>20 (57)</td>
<td>18 (23)</td>
</tr>
</tbody>
</table>

Abbreviations: abs = antibodies; ADEM = acute disseminated encephalomyelitis; AQP4 = aquaporin-4; CIS = clinically isolated syndrome; MOG = myelin oligodendrocyte glycoprotein; MS = multiple sclerosis; NMOSD = neuromyelitis optica spectrum disorder; OCB = IgG bands; ON = optic neuritis; OR = odds ratio with 95% CI; TM = transverse myelitis.

second subgroup consists of children who are older at onset (median 8 years) and predominantly females. ON at onset or later is common, and they are primarily diagnosed with ADEM or CIS but develop a relapsing non-MS course such as MDEM, ADEMON, NMOSD, or recurrent ON with persistent MOG-abs. Furthermore, our results suggest that patients with a recurrent non-MS disease course have higher MOG-ab titers at onset compared with patients with a monophasic course or MS and that persisting MOG-abs are predominantly present in pediatric patients with recurrent non-MS disease course. We conclude that persisting high-titer MOG-abs are associated with a risk of relapse but not the sole presence at onset. Our results also suggest that a cutoff higher than previously anticipated, e.g., ≥1,280, could be used to increase the specificity for a non-MS disease course. However, because this leads to a decrease in sensitivity improvement of MOG-ab testing, using more specific assays as recently published is needed.21 The presence of MOG-abs has also been described in adult patients with MS with a relapsing course involving the brainstem, spinal cord, and poor response to MS therapies. MOG-abs in this subgroup were present over a long time with fluctuation and reappearance of MOG-abs. The authors concluded that patients with MS with MOG-abs represent a distinct phenotype benefiting from different treatment strategies.22 This is in line with biopsy findings in adults with fulminant episodes and MOG-abs revealing a lesion pattern reminiscent of MS type 2 with demyelination and presence of complement activation.23 Our observations are further supported by a recent study involving 50 adults with a history of ON and/or TM: Every fourth positive patient with ON or myelitis had –1). Two children with an MS-like MRI and positive OCB were diagnosed already at age 4 and 8 years. One had an isolated ON and the other patient a relapsing course involving the brainstem, spinal cord, and poor response to MS therapies. MOG-abs in this subgroup were present over a long time with fluctuation and reappearance of MOG-abs. The authors concluded that patients with MS with MOG-abs represent a distinct phenotype benefiting from different treatment strategies.22 This is in line with biopsy findings in adults with fulminant episodes and MOG-abs revealing a lesion pattern reminiscent of MS type 2 with demyelination and presence of complement activation.23 Our observations are further supported by a recent study involving 50 adults with a history of ON and/or TM: Every fourth positive patient with ON or myelitis had –1). Two children with an MS-like MRI and positive OCB were diagnosed already at age 4 and 8 years. One had an isolated ON and the other patient a right-sided hemiplegia. The third patient was 14...
years old with sensory disturbances in his legs due to TM. The 2 younger patients had frequent episodes not involving the brainstem or myelon. All patients are currently treated with MS therapies and MOG-abs dropping to undetectable levels over time. Children with persistent MOG-abs in our study had more episodes with ON compared with children with absent or transient MOG-abs as previously reported.5,6,11,26,27 ON with MOG-abs has distinct features separating it from other diseases such as bilateral involvement, rapid visual impairment combined with good recovery after steroid treatment, and optic disc swelling.26,28–30 Several limitations need to be addressed: MRI studies were performed on scanners with different field strengths, potentially leading to different results. Serologic follow-up was not available from a large part of initially MOG-ab–negative children, as well as from MOG-ab–positive patients without relapses. With a higher number of follow-up serum samples, further conclusions, i.e., regarding the correlation between disease activity and MOG-ab titers, would have been possible. We also

<table>
<thead>
<tr>
<th>MOG-ab titer level at baseline—cutoff values</th>
<th>≥1/160</th>
<th>≥1/320</th>
<th>≥1/640</th>
<th>≥1/1,280</th>
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<tbody>
<tr>
<td>Number, n (%)</td>
<td>65 (31)</td>
<td>59 (28)</td>
<td>52 (25)</td>
<td>41 (19)</td>
</tr>
<tr>
<td>Age, y</td>
<td>6 (0–17)</td>
<td>6 (0–17)</td>
<td>5 (0–17)</td>
<td>5 (0–17)</td>
</tr>
<tr>
<td>Diagnosis at onset, n (%)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>ADEM (60)</td>
<td>34 (57)</td>
<td>33 (55)</td>
<td>30 (50)</td>
<td>27 (45)</td>
</tr>
<tr>
<td>NMO (12)</td>
<td>3 (25)</td>
<td>3 (25)</td>
<td>2 (17)</td>
<td>1 (8)</td>
</tr>
<tr>
<td>CIS (101)</td>
<td>25 (25)</td>
<td>21 (21)</td>
<td>19 (19)</td>
<td>13 (13)</td>
</tr>
<tr>
<td>MS (37)</td>
<td>3 (8)</td>
<td>2 (5)</td>
<td>1 (3)</td>
<td>0 (0)</td>
</tr>
<tr>
<td>Disease course (monophasic, recurrent non-MS, MS) after 2 y, n (%)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Monophasic (98)</td>
<td>40 (42)</td>
<td>36 (37)</td>
<td>33 (34)</td>
<td>25 (26)</td>
</tr>
<tr>
<td>ADEM (48)</td>
<td>22 (48)</td>
<td>21 (46)</td>
<td>20 (43)</td>
<td>19 (39)</td>
</tr>
<tr>
<td>CIS (47)</td>
<td>15 (32)</td>
<td>12 (25)</td>
<td>11 (23)</td>
<td>6 (13)</td>
</tr>
<tr>
<td>NMO (3)</td>
<td>3 (100)</td>
<td>3 (100)</td>
<td>2 (67)</td>
<td>1 (33)</td>
</tr>
<tr>
<td>Recurrent non-MS (35)</td>
<td>22 (63)</td>
<td>21 (60)</td>
<td>18 (51)</td>
<td>16 (46)</td>
</tr>
<tr>
<td>MDEM/ADENON (11)</td>
<td>11 (100)</td>
<td>11 (100)</td>
<td>9 (82)</td>
<td>8 (73)</td>
</tr>
<tr>
<td>NMO (13)</td>
<td>3 (23)</td>
<td>3 (23)</td>
<td>3 (23)</td>
<td>3 (23)</td>
</tr>
<tr>
<td>Recurrent ON (11)</td>
<td>8 (73)</td>
<td>7 (64)</td>
<td>6 (54)</td>
<td>5 (45)</td>
</tr>
<tr>
<td>MS (79)</td>
<td>3 (4)</td>
<td>2 (2)</td>
<td>1 (1)</td>
<td>0 (0)</td>
</tr>
</tbody>
</table>

Prediction of a non-MS disease course (monophasic or recurrent) after 2 y

| Sensitivity (95% CI) | 0.473 (0.390, 0.558) | 0.435 (0.353, 0.521) | 0.389 (0.310, 0.475) | 0.313 (0.240, 0.397) |
| Specificity (95% CI) | 0.962 (0.894, 0.990) | 0.975 (0.912, 0.996) | 0.987 (0.932, 0.999) | 1.0 (0.954, 1.0) |
| ρ | <0.0001 | <0.0001 | <0.0001 | <0.0001 |

Prediction of a recurrent non-MS disease course after 2 y

| Sensitivity (95% CI) | 0.629 (0.463, 0.768) | 0.6 (0.436, 0.744) | 0.514 (0.356, 0.670) | 0.457 (0.305, 0.618) |
| Specificity (95% CI) | 0.754 (0.685, 0.812) | 0.783 (0.716, 0.837) | 0.806 (0.741, 0.857) | 0.857 (0.780, 0.901) |
| ρ | <0.0001 | <0.0001 | <0.0002 | <0.0001 |

Abbreviations: abs = antibodies; ADEM = acute disseminated encephalomyelitis; ADEMON = acute disseminated encephalomyelitis with optic neuritis; CIS = clinically isolated syndrome; MDEM = multiphasic disseminated encephalomyelitis; MOG = myelin oligodendrocyte glycoprotein; MS = multiple sclerosis; NMO = neuromyelitis optica spectrum disorder; OCB = IgG bands; ON = optic neuritis.

*Median (range).

bCalculated using the Fisher exact test.
have to mention that 63 children of our cohort were reported in previous studies, potentially leading to a selection bias. Nevertheless, the proportion of MOG-ab-positive children was similar to other reported cohorts.11,12

Since a previous publication indicated that MOG-ab reactivity in patients with MS might be caused by IgM abs detected by IgG heavy + light chain (H + L) reactive secondary abs, whereas IgG1 reactive secondary abs are more specific,21 we tested a subset of our samples with different secondary abs (table e-5). Although IgM abs were present in some of these samples, all were also IgG1 reactive. This finding is consistent with the work of Waters et al.,21 who showed that almost all IgG1-reactive samples also have MOG-antibody titers ≥1:160 when measured using the (H + L) assay.

MOG-abs are present in a substantial number of very young children with ADS and are associated with a monophasic disease course in the context of declining antibody levels. High and persistent MOG-abs on the other hand are associated with a recurrent disease course other than MS. Collaborative studies are needed to better define this group of diseases with the common denominator MOG-abs.

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AUTHOR CONTRIBUTIONS
Eva-Maria Hennes: acquisition of data, study supervision, and analysis and interpretation of data. Matthias Baumann: acquisition of data, analysis and interpretation of data, and critical revision of the manuscript for intellectual content. Kathrin Schanda: acquisition of data and analysis and interpretation of data. Banu Anlar: Barbara Bajer-Kornek, Astrid Bläshek, Sigrid Blatter-Inselder, Katharina Diepold, Astrid Eisenkölbl, Thaddäus Gotzwald, Georgi Kuchukidze, Ursula Gruber-Steinmayr, Martin Häusler, Romana Höfberger, Michael Karentorf, Andrea Klein, Johannes Koch, Verena Krause, Christian Lechner, Steffen Leiz, Frank Leypoldt, Simon Mader, Klaus Marquard, Imke Poggenburg, Daniela Pohl, Martin Prinzhorn, Markus Rauchzehnauner, Mareike Schimmel, Charloste Theis, Daniel Tibussek, Silvia Vicker, and Carolin Zeches: acquisition of data and critical revision of the manuscript for intellectual content. Thomas Berger: acquisition of data, analysis and interpretation of data, and critical revision of the manuscript for intellectual content. Markus Rostasy: acquisition of data, analysis and interpretation of data, and study supervision. Kevin Rostasy: study concept and design, acquisition of data, analysis and interpretation of data, and study supervision.

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