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Partial acute transverse myelitis is a predictor of multiple sclerosis in children

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Abstract
Background: Acute transverse myelitis (ATM) in children is a rare and often severe disease for which there are few known prognostic factors, particularly the subsequent risk of multiple sclerosis (MS) diagnosis.
Objectives: To determine the clinical course and prognostic factors after a first episode of ATM in children.
Methods: Thirty children below 16 years of age diagnosed with a first neurological episode of ATM were included retrospectively. Clinical evaluation, treatment, laboratory, and MRI data were collected.
Results: Median age at onset was 11 years (range 3–15 years). Follow-up data were available for a median of 4 years (range 0.5–16.7 years). Five patients subsequently had a diagnosis of MS (17%), which was associated with acute partial transverse myelitis (odds ratio 5; 95% confidence interval 2.3–11), with a 60% probability of having a relapse at five years (p < 0.01). The 2011 Verhey criteria correctly identified MS in children with the highest specificity (96%) and sensitivity (80%).
Conclusion: Acute partial transverse myelitis and brain MRI abnormalities at initial presentation are significantly predictive of a subsequent diagnosis of MS in children with ATM. These findings suggest that closer brain MRI monitoring after acute partial transverse myelitis might make the earlier introduction of disease-modifying therapies possible.

Keywords
Multiple sclerosis, acute transverse myelitis, pediatric, magnetic resonance imaging

Introduction
Acute transverse myelitis (ATM) is a rare inflammatory process affecting the spinal cord, which manifests as sensory, motor or autonomic dysfunction, with an incidence of 1.71 to 2 per million children.¹ ² Precise diagnostic criteria for idiopathic and non-idiopathic ATM have been proposed by the Transverse Myelitis Consortium Working Group.³

The American Academy of Neurology (AAN) published evidence-based ATM management guidelines in 2011, focusing on two clinical settings: acute complete transverse myelitis (ACTM), defined as acute or subacute inflammation of the spinal cord causing moderate or severe loss of function, and acute partial transverse myelitis (APTM), characterized by incomplete or patchy involvement of at least one spinal segment, with mild to moderate weakness, asymmetric or dissociated sensory symptoms and, occasionally, bladder involvement.⁴ In adult patients, this distinction is considered useful for determining ATM etiology and relapse risk.
We hypothesized that this might also be the case in children. We investigated the prognostic factors of childhood ATM, focusing particularly on the risk of multiple sclerosis (MS) diagnosis and disability status at last follow-up.

Method

Patients

Inclusion criteria. All children under the age of 16 years evaluated at Montpellier University Hospital (CHRU de Montpellier), France, from January 1994 to December 2009, with a diagnosis of ATM as a first neurological episode, were included. ATM diagnosis was based on the 2002 criteria. If spinal magnetic resonance imaging (MRI) showed an appropriately located high signal-intensity lesion on T2-weighted sequences but no clear cut enhancement of the abnormality following gadolinium administration and no CSF abnormality, patients were considered to have ‘possible ATM’ and were included.

Exclusion criteria. Patients with a previous neurological history, systemic or metabolic disease, history of previous spinal radiation, extra-axial spine compression or thrombosis of the anterior spinal artery on MRI were excluded.

Standard protocol approvals, registrations, and patient consent

The parents of the patients gave written informed consent to care. The procedures described herein are all part of routine care on our hospital. The study has been submitted to the ethics committee of Montpellier University Hospital: this observational and retrospective study did not require approval of the institutional review board and no informed written consent was required, in line with French law.

Data collection

All the information collected was coded to protect confidentiality.

Clinical data. Retrospective clinical data during acute phase were collected from our hospital database in a standardized manner: demographic, personal, and familial history of MS and other autoimmune disease, factors preceding ATM, such as vaccination, trauma, or infection. The acute illness description included first symptoms, time to diagnosis and maximal clinical expression, and plateau duration. Patient management, particularly the need of bladder catheterization, and the type of treatment were also recorded.

Walking ability during the acute phase was assessed with a modified version of the Hughes Functional Disability Scale (HFDS) previously used in Guillain–Barré syndrome, and in the largest published pediatric ATM series:

\[
0 = \text{normal}, \ 1 = \text{minor symptoms, fully capable of manual work}, \ 2 = \text{able to walk more than 30 feet without assistance}, \ 3 = \text{able to walk more than 30 feet with assistance}, \ 4 = \text{bed-bound/wheelchair-bound}, \ 5 = \text{requiring assisted ventilation}, \ 6 = \text{dead}.
\]

Laboratory and MRI data. We collected the results of cerebrospinal fluid (CSF) tests, including white blood cell (WBC) count, biochemistry findings, and oligoclonal band detection. Serum samples were tested for 19 infectious pathogens, with convalescent titers for the same pathogens determined within six weeks of the acute phase, when possible. An infectious disease was defined by two recorded oral temperatures above 38.5°C (101.3°F), two recorded serum WBC elevations above 11,000 cells/mm³, or positive PCR or IgM converting to IgG against a specific pathogen.

Magnetic resonance images were acquired on 1.5 T scanners. MRI data included scans of the spine and brain at the same time during the acute phase and follow-up: localization, extent and type of spine lesions, gadolinium enhancement on T1-weighted images (T1W). Patients with lesions at least three segments long were classified as having ‘longitudinally extending transverse myelitis’ (LETM). MRI brain images were examined for the presence of T2-weighted (T2W) lesions (supratentorial, infratentorial, basal ganglia/thalamus) and were classified according to: (1) the 2010 McDonald dissemination in space and time criteria; (2) the 2011 Verhey criteria; (3) the KIDMUS criteria; (4) the CHAMPS criteria. Criteria definitions are shown in Table 1.

All baseline images were reviewed retrospectively, with a standardized MRI record form, by an experienced neuroradiologist (NL) and neuropediatrician (FR) blinded to clinical symptoms, disease progression, and initial MRI analysis, to obtain a consensus. Visual evoked potentials (VEPs) were also considered, when performed.

Outcome data. Outcome data were collected from our hospital database for patients having ongoing care provided by our department, or from physicians caring for other children in the community. Outcomes included subsequent diagnosis of MS and Kurtzke Expanded Disability Status Scale (EDSS) score at last follow-up, focusing on scores ≥4, defined as limited walking ability but able to walk for more than 500 m without assistance or resting.

Definitions and classification

Patients with incomplete or patchy involvement of at least one spinal segment, with mild to moderate weakness, asymmetric or dissociated sensory symptoms and, occasionally, bladder involvement were defined as having APTM. Patients with symmetric, moderate or severe loss of function due to a spinal cord lesion were considered to have ACTM.
### Table 1. Test properties of brain MRI criteria for predicting conversion to multiple sclerosis.

<table>
<thead>
<tr>
<th>Definition</th>
<th>McDonald (2010)</th>
<th>Verhey</th>
<th>KIDMUS</th>
<th>CHAMPS</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dissemination in space criteria: ≥1 T2 hyperintense lesion in ≥2 of the following regions: periventricular, juxtacortical, infratentorial, spinal cord (symptomatic spinal cord lesion excluded)</td>
<td>85–100</td>
<td>20</td>
<td>62–84</td>
<td>80</td>
</tr>
<tr>
<td>Dissemination in time criteria: Simultaneous presence of asymptomatic gadolinium-enhancing and non-enhancing lesions at any time</td>
<td>93–100</td>
<td>92</td>
<td>84–93</td>
<td>96</td>
</tr>
<tr>
<td>Both dissemination in space and dissemination in time</td>
<td>58–100</td>
<td>96</td>
<td>82–94</td>
<td>96</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Ref.</th>
<th>Sensitivity</th>
<th>Specificity</th>
<th>Positive predictive value</th>
<th>Negative predictive value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Our series</td>
<td>80</td>
<td>83.3</td>
<td>50</td>
<td>95.2</td>
</tr>
<tr>
<td>Our series</td>
<td>85–100</td>
<td>66–80</td>
<td>36</td>
<td>100</td>
</tr>
<tr>
<td>Our series</td>
<td>20</td>
<td>88</td>
<td>25</td>
<td>84.6</td>
</tr>
<tr>
<td>Our series</td>
<td>93–100</td>
<td>84–88</td>
<td>55</td>
<td>100</td>
</tr>
<tr>
<td>Our series</td>
<td>20</td>
<td>92</td>
<td>50</td>
<td>85.7</td>
</tr>
<tr>
<td>Our series</td>
<td>58–100</td>
<td>86–100</td>
<td>59</td>
<td>96</td>
</tr>
<tr>
<td>Our series</td>
<td>80</td>
<td>96</td>
<td>80</td>
<td>76</td>
</tr>
<tr>
<td>Our series</td>
<td>62–84</td>
<td>84–93</td>
<td>80</td>
<td>59</td>
</tr>
<tr>
<td>Our series</td>
<td>60</td>
<td>95.8</td>
<td>75</td>
<td>96</td>
</tr>
<tr>
<td>Our series</td>
<td>8–57</td>
<td>87.5</td>
<td>92</td>
<td>58</td>
</tr>
<tr>
<td>Our series</td>
<td>80</td>
<td>95–100</td>
<td>78</td>
<td>57.1</td>
</tr>
<tr>
<td>Our series</td>
<td>78</td>
<td>63–87</td>
<td>95.5</td>
<td>84</td>
</tr>
<tr>
<td>A12b</td>
<td>58</td>
<td>69</td>
<td>84</td>
<td></td>
</tr>
</tbody>
</table>

Values are percentages. All four criteria correctly identified patients at risk of transition to MS (p < 0.05). P = Pediatric study A = Adult study.

### General overview

**Clinical findings** (no clear arterial distribution clinical signs before ATM onset was described in 17% of cases (5/30). In 60% (18/30) of cases, neurologic manifestations developed after an infection occurring a median of 6 days (range 2–28 days) before the onset of neurologic symptoms: isolated fever (7/18), encephalopathy (6/18), meningoencephalitis (1/18), pyogenic meningitis (1/18), and 6/20 (30%) underwent neurologic testing after one day to six weeks. Minor trauma (fall or twist) up to one day before ATM onset was described in 17% of cases (5/30). In total, 30 patients with acute transverse myelitis before the age of 16 years were included in the study. The mean follow-up of 5.1 ± 4.4 years (median: 3.5 years) was 11.3 years. At the latest follow-up, 17% (5/30) had ATMs, 17% (5/30) had MS, 13% (4/30) had recurrent ATM, and 6/30 (20%) had encephalopathy. The main clinical, CSF, and MRI characteristics are shown in Table 2.

**Diagnostic criteria** (no clear arterial distribution clinical signs before ATM onset was described in 17% of cases (5/30). ATM was classified as: isolated ATM with no clinical brain disease and normal brain MRI, clinically isolated syndrome (CIS), i.e. clinically isolated ATM clinically isolated syndrome (CIS) with no clinical brain disease and normal brain MRI, clinically isolated syndrome (CIS) with encephalopathy, polyfocal CIS, i.e. multifocal neurologic deficit including ATM, or MS fulfilling the 2010 McDonald criteria.

### Results

**Statistical analysis**

Descriptive data were compared in Χ² tests or Fisher’s exact tests for categorical variables and t-tests or Wilcoxon tests for continuous measures. Kaplan–Meier curves were plotted and the significance of differences was assessed in log-rank tests. Statistical significance was assessed in a value of less than 0.05. We were unable to conduct multivariate analysis due to the small size of our population.
Table 2. Main clinical, CSF, and MRI characteristics of the patients during the acute phase.

<table>
<thead>
<tr>
<th>Clinical data</th>
<th>Total</th>
<th>Isolated ATM</th>
<th>CIS</th>
<th>MS</th>
<th>ADEM</th>
<th>Other*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Preceding infection (%)</td>
<td>60</td>
<td>53</td>
<td>100</td>
<td>20</td>
<td>75</td>
<td>100</td>
</tr>
<tr>
<td>Median duration to maximal functional symptoms [d] (range)</td>
<td>5 (1–17)</td>
<td>3 (1–12)</td>
<td>6 (3–7)</td>
<td>5 (4–5)</td>
<td>6 (2–12)</td>
<td>17</td>
</tr>
<tr>
<td>Pain (%)</td>
<td>67</td>
<td>71</td>
<td>100</td>
<td>40</td>
<td>50</td>
<td>100</td>
</tr>
<tr>
<td>Fever (%)</td>
<td>40</td>
<td>40</td>
<td>60</td>
<td>20</td>
<td>25</td>
<td>100</td>
</tr>
<tr>
<td>Complete paraplegia (%)</td>
<td>46</td>
<td>53</td>
<td>40</td>
<td>20</td>
<td>75</td>
<td>100</td>
</tr>
<tr>
<td>Pyramidal signs (%)</td>
<td>50</td>
<td>47</td>
<td>40</td>
<td>75</td>
<td>50</td>
<td>0</td>
</tr>
<tr>
<td>Sensory loss (%)</td>
<td>60</td>
<td>79</td>
<td>40</td>
<td>60</td>
<td>25</td>
<td>100</td>
</tr>
<tr>
<td>Sphincter deficit (%)</td>
<td>77</td>
<td>47</td>
<td>20</td>
<td>20</td>
<td>25</td>
<td>0</td>
</tr>
<tr>
<td>Bladder catheterization (%)</td>
<td>33</td>
<td>21</td>
<td>40</td>
<td>100</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Median HFDS score (range)</td>
<td>3.5 (1–4)</td>
<td>4 (1–4)</td>
<td>3 (2–4)</td>
<td>3 (2–4)</td>
<td>4 (3–4)</td>
<td>4</td>
</tr>
<tr>
<td>Median CSF WBC count [cells/mm$^3$] (range)</td>
<td>15.5 (0–700)</td>
<td>10 (0–700)</td>
<td>30 (2–76)</td>
<td>11 (2–64)</td>
<td>200 (30–530)</td>
<td>20</td>
</tr>
<tr>
<td>Median CSF protein concentration [g/l] (range)</td>
<td>0.45 (0.15–1.79)</td>
<td>0.39 (0.21–0.89)</td>
<td>0.38 (0.15–0.79)</td>
<td>0.44 (0.21–0.67)</td>
<td>0.69 (0.42–1.79)</td>
<td>0.58</td>
</tr>
<tr>
<td>Median lesion length on spinal MRI [segment] (range)</td>
<td>5 (0–15)</td>
<td>3 (0–15)</td>
<td>10 (4–16)</td>
<td>4 (0–9)</td>
<td>6 (0–7)</td>
<td>20</td>
</tr>
<tr>
<td>Abnormal brain MRI (%)</td>
<td>45</td>
<td>0</td>
<td>100</td>
<td>80</td>
<td>100</td>
<td>0</td>
</tr>
</tbody>
</table>

*Final diagnosis of connective tissue disorder; †Significantly higher in MS than in the non-MS population; ‡Significantly lower in MS than in the non-MS population. ATM = acute transverse myelitis; APTM = acute partial transverse myelitis; CIS = clinically isolated syndrome; MS = multiple sclerosis; ADEM = acute disseminated encephalomyelitis; CSF = cerebrospinal fluid; WBC = white blood cell; HFDS = Hughes Functional Disease Scale.

deficit consistent with thrombosis of the anterior spinal artery, time to reach maximal severity > 4 hours from onset) and spinal MRI findings in these five patients were not consistent with infarction.

The main clinical characteristics are shown in Table 2. Acute partial transverse myelitis was present in 33% of the patients (10/30). By definition, the five patients initially diagnosed with ADEM had encephalopathy during the acute phase. 13

**CSF and MRI characteristics.** Cerebrospinal fluid (CSF) was obtained by lumbar puncture, 1 to 13 days after symptom onset (median 2.5 days). Abnormalities were found in 17 patients (57%, see Table 2). CSF oligoclonal bands were detected in 4 of 16 patients tested (3 MS and 1 post-infectious polyfocal CIS) and were not significantly associated with MS.

All patients underwent a first cerebral and spinal MRI between 1 and 11 days after symptom onset (median: 4 days) and images were available for 29 patients. Spinal MRI was abnormal in 93% of cases (27/29): T2-weighted (T2W) signal abnormalities were identified at cervical level in 55% patients (16/29), thoracic level in 72% (21/29) and lumbosacral level in 24% (7/29). Multifocal lesions were detected in 26% of patients (8/29) and hypointense lesions on T1-weighted (T1W) sequences were found in two patients. LETM was found in 20 of 29 patients (69%). Total median segment length was 5 segments (range 0–15 segments). Lesion count did not differ significantly between patients with and without MS ($p = 0.283$). Gadolinium infusion demonstrated lesion enhancement on T1W images in 31% (9/29) of cases. Both patients who had a first normal spinal MRI showed lesions on T2W sequences without gadolinium enhancement on a second scan performed 5 and 7 days later.

Cerebral MRI evidence of white matter lesions was obtained in 45% (13/29) of cases: the hypersignal areas on T2W images were supratentorial in 38% (11/29) of cases, infratentorial in 31% (9/29), and in the basal ganglia/thalamus in 28% (8/29) of cases. One patient (final diagnosis: MS) had a T2W hypersignal on optic nerves. The test properties of the 2010 McDonald, Verhey, KIDMUS, and CHAMPS criteria are shown in Table 1.

**Visual evoked potentials (VEP) and immunochemistry.** VEP was carried out during the acute phase in 60% of patients (18/30) and optic neuritis was found in 13% of cases (4/30). Serum anti-aquaporin 4 IgG was absent in all four children.
tested: only 1 of 4 had optic neuritis, with LETM and a subsequent diagnosis of MS. None of the patients presented serologic evidence of connective tissue disease at presentation but, during follow-up, one patient displayed antinuclear antibodies, lupus anticoagulant and ANCA, suggestive of this condition.

**Type of treatment.** All patients were prescribed treatment at ATM presentation: 87% (26/30) received high-dose IV methylprednisolone, 50% (15/30) received IV immunoglobulin, and both were prescribed in 37% of cases (11/30). Mean interval between treatment beginning and MRI was of 0.5 days (median: 0 days, range: 4 days before MRI to 5 days after first MRI). We found no correlation between type of treatment and outcome.

**Prognosis factors**

Follow-up duration was 5.1 ± 4.4 years (range: 0.5–16.7) and exceeded one year in more than 96% of cases (29/30). Most (80%) patients (24/30) were able to walk independently within one month of onset. The other six patients who were unable to walk independently within one month of onset had a median HFDS score of 4 during acute phase (range: 3–4), a median plateau duration phase of 10 days (range: 5–20 days), and a median lesion length of 6 segments on spinal MRI (range: 2–15 segments). Outcome was not related to seasonality, sex, age at onset, time to maximal symptom expression, fever, pain, pyramidal signs at the time of presentation, CSF findings or spinal MRI characteristics during acute phase.

**Diagnosis of MS.** Acute partial transverse myelitis at initial presentation was a significant prognostic factor for subsequent MS diagnosis (odds ratio [OR] 5, 95% confidence interval [CI] 2.3–1, \( p < 0.01 \)) with a 40% probability of being relapse-free at five years \( (p < 0.01, \text{see Figure 2}).\) Higher acute-phase HDFS score (3 or 4) was associated with significant protection against diagnosis of MS (OR 0.048, 95%CI 0.01–0.54, \( p < 0.05 \)).

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*Figure 1.* Patient diagnosis flow chart. ATM = acute transverse myelitis; ‘possible’ ATM = clinical ATM with T2 hyperintensity on spinal MRI without gadolinium enhancement, and normal CSF;³ CIS = clinically isolated syndrome; monofocal CIS = ATM with abnormal brain MRI, without encephalopathy; polyfocal CIS = multifocal neurological deficit including ATM, without encephalopathy; ADEM = acute demyelinating encephalomyelitis; MS = multiple sclerosis.
The 2010 McDonald dissemination in space, Verhey, KIDMUS, and CHAMPS criteria correctly identified patients at risk of MS diagnosis ($p < 0.05$; see Table 1).

**Disability scale score at last follow-up assessment (EDSS).** EDSS score at last follow-up was significantly higher in patients taking more than one month to walk independently (mean of 4.3 vs. 0.9, $p < 0.01$) and in cases of complete paraplegia during acute phase (mean 2.3 vs. 0.8, $p < 0.05$). The need for urinary catheterization during acute phase was significantly associated with a final EDSS score $\geq 4$ (OR 2.5, 95% CI 1.5–4, $p < 0.05$). Patients with idiopathic ATM were more likely to have an EDSS score $\geq 4$ at one year (OR 24, 95% CI 1.5–395, $p < 0.05$), but this difference was not significant at the end of follow-up.

**Discussion**

This retrospective series of children below the age of 16 years with single episodes of ATM at a single academic center included 30 patients over a 15-year period. Previous series specifically studying pediatric ATM concerned eight to 47 patients included over four to 25 years. Apart from one, all were monocentric and retrospective, and not all used the 2002 diagnostic criteria. Other larger studies of pediatric cases of acute CNS inflammatory demyelination did not include details of clinical symptoms at presentation. It is difficult to compare our results with those of previously published series, as the 2002 criteria (including markers of spinal inflammation) were not used in five of these studies. The clinical presentation and course of ATM in our patients was similar to those in previous pediatric series, in terms of age at presentation, initial course and preceding infection (see Table 3). Surprisingly, none of our patients was under three years old, contrasting with the two largest published series, which described a bimodal incidence distribution, with peaks before the age of five years and during adolescence. We also found a bimodal distribution, but with peaks in the 3–5 years and 10–15 years age groups. Unlike some other studies, we found no relationship between young age at presentation and prognosis. Sex did not influence outcome. As previously reported, none of our patients was vaccinated or had an allergy shot within 30 days of the first symptoms of ATM, by contrast to a previous center-based analysis of 47 cases.

Idiopathic ATM had the worst prognosis at one year (EDSS $\geq 4$), but this difference was not significant at the end of follow-up. In a study of idiopathic ATM in adults with a mean follow-up period of 3.6 ± 3.1 years, 30% of patients had a poor outcome, defined as death or a loss of walking ability. Consistent with previous reports, we observed potential functional recovery after one year. It remains unclear whether this is due to primary neurological recovery, reeducation or both.

The main finding of this study was the five times higher risk of subsequent MS diagnosis in children with acute partial transverse myelitis at initial presentation. To our knowledge, this is the first study to focus on APTM and MS risk in children. Since 1992, ATM has been divided into two subgroups by several authors: acute complete transverse myelitis (ACTM) and acute partial transverse myelitis (APTM), which differ in terms of clinical presentation, prognosis and course. This clinical dichotomy was highlighted in 2011, in the American Academy of Neurology (AAN) evidence-based guidelines. The 2002 criteria classified patients into relatively homogeneous groups in terms of clinical and MRI data, but outcome remained unpredictable (except for ‘spinal shock’ with complete paraplegia). It should also be noted that all four of our initial clinical scenarios wound up including MS cases. It was therefore necessary to identify other prognostic factors in ATM. In the AAN evidence-based guidelines, a review of two studies of APTM with normal brain MRI in adults reported subsequent diagnosis of MS in 10.3% of cases (95% CI 4.1–23.6), whereas two studies of adults with ACTM and normal brain MRI suggested a significantly lower subsequent diagnosis of MS (0% to 2%), over about five years of follow-up, as in our study. For adults with APTM and cerebral lesions on MRI, the subsequent diagnosis of MS was significantly higher, at 59 to 90%, by comparison, subsequent MS diagnosis in our pediatric series was of 20% for APTM with normal brain MRI, 66% for APTM with abnormal brain MRI, and 0% for ACTM. Our results suggest that, as in adults, cerebral MRI status at presentation is the most reliable indicator of the risk of subsequent MS.
| Reference       | Country | Year of publication | Number of patients | Study period | Mean follow-up, y | Sex, M:F | Mean age at onset, y (range) | Age at onset <3 y, % | ATM following infection, % | Mean time from infection to onset, d | Mean time to maximal deficit | Pain (any site), % | Backache, % | Fever, % | Complete paraplegia, % | Acute sphincter dysfunction, % | Acute sensory level | Abnormal CSF findings, % (n) | Abnormal spinal cord MRI findings, % (n) | Death, n | Complete recovery, % | Multiple sclerosis, n |
|-----------------|---------|---------------------|--------------------|--------------|-------------------|----------|----------------------------|---------------------|--------------------------|-----------------------------------|-----------------|----------------|-------------|----------------|-----------------------------|------------------|-----------------------------|-------------------------------|---------|------------------|------------------|
| Thomas et al.14 | Canada  | 2012                | 38                 | 1999–2006    | 3.2               | 10.9 (0.5–17) | 5                   | 60                  | 10                        | 8.8                          | 11              | 67              | 60          | 50             | 77                          | 3                | 63                          | 86 (19/30)                   | 93 (27/29) | 48               | 5                |
| De Goede et al.2 | UK      | 2010                | 41                 | 2002–2004    | 0.5               | 8.7 (0–15)       | 20                  | 60                  | 12                        | 1.66                         | 5               | 68              | 34          | 63             | 47                          | 7                | 63                          | 68 (21/31)                   | 69 (27/39) | 52               | 0                |
| Pidcock et al.6  | USA     | 2007                | 8                  | 2000–2004    | 8                 | 8.3 (0–14)       | 38                  | 63                  | 13                        | 0.85                         | 7               | 53              | 59          | 89             | 63                          | 10               | 63                          | 69 (17/36)                   | 69 (27/39) | 59               | 3                |
| Defresne et al.15 | France  | 2003                | 24                 | 1965–1995    | 7.3               | 7 (0–14)         | 12.5                | 100                 | 10                        | 2.5                          | 5               | 83              | 53          | 89             | 85                          | 85               | 85                          | 71 (24/34)                   | 91 (19/21) | 52               | 0                |
| Knebusch et al.17 | Germany | 1998                | 8                  | 1993–1996    | 1.66              | 1.66 (0–15)      | 0                   | 100                 | 10                        | 5                            | 11              | 83              | 80          | 65             | 85                          | 60               | 90                          | 62 (15/24)                   | 60 (12/20) | 75               | 0                |
| Dunne et al.16   | Australia | 1986                | 21                 | 1966–1983    | 0.9               | 0.9 (0.5–15)     | 0                   | 100                 | 10                        | 10                           | 5               | 53              | 80          | 65             | 85                          | 5                | 65                          | 71 (24/34)                   | 50 (4/8) | 52               | 0                |
| Paine and Byers18 | USA     | 1953                | 25                 | 1929–1952    | 0.5               | 0.5 (0–15)       | 1                   | 100                 | 10                        | –                            | –               | 53              | –           | 53             | 53                          | 53               | 53                          | 53 (36/38)                   | 53 (36/38) | 0                | 0                |
diagnosis in children with APTM. Yearly cerebral MRI for at least five years, with clinical assessments, has been suggested, to monitor adult patients with APTM for MS diagnosis.22 Our results support this suggestion, which also appears reasonable for children, to optimize the early diagnosis of MS.

We compared the performances of the 2010 McDonald, Verhey, KIDMUS, and CHAMPS criteria for predicting subsequent diagnosis of MS (see Table 1). Table 1 also shows previous published series studying these criteria, but it is to note that none of them focused specifically on acute transverse myelitis.9,11,29-32 In our series, all criteria had low sensitivity, but high specificity, the highest values being those for the Verhey criteria. These properties have already been reported for children, particularly those older than 10, for KIDMUS criteria.9,10 It has also been shown that the 2010 McDonald had high sensitivity and specificity (100% and 86%, respectively) or children older than 11 years with non–ADEM presentations.39 The greatest advantage of the Verhey, 2010 McDonald, and KIDMUS scales was their high negative predictive value. Moreover, the low positive predictive value (PPV) of the 2010 McDonald criteria (50%) highlighted the need for specific pediatric scales, such as the Verhey criteria, with a higher PPV (80%). This is the first time that the CHAMPS criteria have been studied in a pediatric population, but they were not found to be particularly useful, particularly given their very low PPV.

Gadolinium infusion demonstrated lesion enhancement on T1W images in 31% of spinal MRI scans and may explain the low McDonald Dissemination in Time criteria in our cohort. Gadolinium enhancement in pediatric ATM series is of 24 to 74%,6,14 leading to suggest that the 2002 consortium guidelines may underdiagnose pediatric ATM, but this needs further studies.

One of the five patients manifesting with an ADEM phenotype at onset had two relapses, without encephalopathy, during follow-up and was diagnosed with MS. ADEM-like episode as a first event of MS is reported in 2 to 18% of cases in published pediatric series of patients with demyelination.9,19,33

As reported for both adult and pediatric ATM, a ‘spinal shock’ presentation (complete paraplegia, hypotonia, and areflexia) was associated with a poor EDSS score at the end of follow-up.2,15,21,34 However, unlike other studies, we found no relationship between poor prognosis and higher sensitive level,5 pyramidal syndrome,20 or supraspinal symptoms.15 We confirmed that taking more than one month to walk independently was associated with a poor EDSS score at the end of follow-up.15

We found no correlation between spinal lesion length and subsequent diagnosis of MS, probably due to our small number of MS cases, or EDSS score at the end of follow-up. In adult MS patients, the risk of such diagnosis is higher in patients with spinal lesions less than two sections long23, but this result has never been confirmed in children.14,15,35,36 We found no association between oligoclonal band detection and MS in our series, probably due to the small number of patients tested. Unlike Pidcock et al.,6 we found no relationship between poor prognosis and lower lesion level and T1 hypointensity on spinal MRI. Most children had LETM, as previously reported,6,36 and none had neuromyelitis optica according to the Wingerchuk criteria.7 Neuromyelitis optica antibody tests were negative in all four children studied.

**Conclusion**

Our single-center analysis of 30 pediatric cases of acute transverse myelitis highlighted the significantly higher risk of multiple sclerosis diagnosis in cases of acute partial transverse myelitis, as previously reported for adults.4 This simple clinical dichotomy (mild to moderate weakness, asymmetric or dissociated sensory symptoms), which can be assessed at the patient’s bedside during the acute phase, provided valuable clinical decision support. Together with brain MRI data, it could help to identify children at higher risk of subsequent diagnostic of multiple sclerosis, leading to closer brain MRI monitoring or to early disease-modifying treatment.

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**Conflict of interest**

The authors declare that there is no conflict of interest.

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**References**