Increased frequency of anti-Ma2 encephalitis associated with immune checkpoint inhibitors

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Neurology Neuroimmunol Neuroinflamm 2019;6:e604. doi:10.1212/NXI.0000000000000604

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Abstract

Objective
To report the induction of anti–Ma2 antibody–associated paraneoplastic neurologic syndrome (Ma2-PNS) in 6 patients after treatment with immune checkpoint inhibitors (ICIs). We also analyzed (1) patient clinical features compared with a cohort of 44 patients who developed Ma2-PNS without receiving ICI treatment and (2) the frequency of neuronal antibody detection before and after ICI implementation.

Methods
Retrospective nationwide study of all patients with Ma2-PNS developed during ICI treatment between 2017 and 2018.

Results
Our series of patients included 5 men and 1 woman (median age, 63 years). The patients were receiving nivolumab (n = 3), pembrolizumab (n = 2), or a combination of nivolumab and ipilimumab (n = 1) for treatment of neoplasms that included lung (n = 4) and kidney (n = 1) cancers and pleural mesothelioma (n = 1). Clinical syndromes comprised a combination of limbic encephalitis and diencephalitis (n = 3), isolated limbic encephalitis (n = 2), and a syndrome characterized by ophthalmoplegia and head drop (n = 1). No significant clinical difference was observed between our 6 patients and the overall cohort of Ma2-PNS cases. Post-ICI Ma2-PNS accounted for 35% of the total 17 Ma2-PNS diagnosed in our center over the 2017–2018 biennium. Eight cases had been detected in the preceding biennium 2015–2016, corresponding to a 112% increase of Ma2-PNS frequency since the implementation of ICIs in France. Despite ICI withdrawal and immunotherapy, 4/6 patients died, and the remaining 2 showed a moderate to severe disability.

Conclusions
We show a clear association between ICI use and increased diagnosis of Ma2-PNS. Physicians need to be aware that ICIs can trigger Ma2-PNS because clinical presentation can be challenging.

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Go to Neurology.org/NN for full disclosures. Funding information is provided at the end of the article.

Data access, responsibility, and analysis: The corresponding author had full access to all the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis.

The Article Processing Charge was funded by Institut Neuromyogène INSERM U1217 CNRS S310.

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Therapy with monoclonal antibodies (Abs) targeting immune checkpoints, including cytotoxic T lymphocyte–associated antigen 4 (CTLA-4), the programmed death-1 receptor (PD-1), and its ligand PD-L1, has led to a paradigm shift in the treatment of numerous types of cancer. Their unprecedented results in controlling tumors at a metastatic stage have come at the expense of an increased risk of developing immune-related adverse events (irAEs), including severe neurologic complications. Because an increasing number of patients will be exposed to immune checkpoint inhibitors (ICIs) in the forthcoming future, it is crucial to identify the main features of neurologic irAEs.

PNS with Abs targeting the intracellular Ma2 antigen characterizes a peculiar form of encephalitis with prominent involvement of limbic, brainstem, and diencephalic structures, usually in association with testicular or lung cancer. Atypical manifestations including narcolepsy-cataplexy, weight gain, sexual dysfunction, and motor neuron syndrome were described and account for the difficulty in diagnosing anti-Ma2 antibody–associated PNS (Ma2-PNS).

We herein report 6 patients who developed autoimmune encephalitis with anti-Ma2 Abs during treatment with ICIs. To assess the relevance of our findings, we analyzed (1) their clinical features compared with a cohort of 44 patients who had developed Ma2-PNS without receiving any ICI and (2) the impact of ICI use on the frequency of Ma2 detection in a national reference center.

Methods

Patient selection

This is a retrospective study of all patients with anti-Ma2-PNS observed after treatment with ICIs and diagnosed at the French National Reference Center for Paraneoplastic Neurological Syndromes in Lyon, France, between January 1, 2017, and December 31, 2018. All patients underwent a comprehensive laboratory examination for suspected PNS as recommended, including an initial assessment with immunohistochemistry on rat brain sections, followed by a second confirmatory test represented by dot blot analysis on recombinant proteins (Euroimmun, Lübeck, Germany, and/or RAVO Diagnostika, Freiburg, Germany) and/or cell-based assays (CBAs) (in-house techniques) for the presence of onconeural Abs. We systematically tested: anti-Hu, Yo, CV2/CrMP5, Ri, Ma2, amphiphysin, GAD65, AK5, NMDA receptor (NMDAR), AMPAR, GABAAR, GABABR, IgLON5, CASPR2, LGI1, and DPPX. Anti-Ma2 specificities were confirmed using in-house CBA and commercial dot blots (Euroimmun, Lubeck, Germany). Clinical and ancillary data were obtained by telephone or email at the time of diagnosis, based on the biological sample, and at least twice a year to assess clinical evolution. Immunotherapy treatment modalities and oncologic therapy were recorded. Outcomes were assessed using the modified Ranking Scale (mRS). The scale ranges from 0 (no symptoms) to 6 (death).

Clinical comparison between ICI-induced Ma2-PNS vs classic Ma2-PNS

Demographic and clinical features of patients with Ma2-PNS triggered by ICIs were compared with those of the overall cohort of patients with Ma2-PNS unrelated to ICI treatment diagnosed in our center between 2002 and 2018 (n = 44).

Frequency of ICI-related PNS

To assess the impact of ICI use on the development of PNS at a national level, we compared the frequency of the different Ab detections in our Reference Center in the biennium 2017–2018 to the biennium 2015–2016 when the use of ICIs in France was at its starting point. The proportion of Ab-positive cases that developed after ICI use was calculated for each Ab specificity.

Statistical analysis

Descriptive analysis is presented as frequencies and percentages for categorical variables and as the median and range for continuous variables. Categorical data were analyzed with the Fisher exact test (2 tailed) and numerical data with the Mann-Whitney U test. Statistical analyses were performed using IBM SPSS Statistics Software (Version 25.0; IBM Corp, Armonk, NY). p Values <0.05 were considered significant.

Standard protocol approvals, registrations, and patient consents

Written consent was obtained from all patients, and the study was approved by the Institutional Review Board of the University Claude Bernard Lyon 1 and Hospices Civils de Lyon.

Data availability

Data reported in this manuscript are available within the article or its supplementary materials. More information regarding the data is available from the corresponding author on reasonable request.
Results

Patients with ICI-associated anti-Ma2 syndromes

Between 2002 and 2018, we identified 50 patients with Ma2-PNS in our center, 6 of which developed the syndrome after ICI treatment in the biennium 2017–2018. All the information on clinical and paraclinical results, together with associated treatments and outcomes of these 6 patients, is summarized in the table. Most of the patients were male (83%), with a median age of 63 years (range: 47–79 years). All were Caucasians. Four of them had an associated non–small-cell lung cancer, 1 a pleural mesothelioma, and the last one a renal clear cell carcinoma. At the time of ICI introduction—a median of 6.5 months (range: 0.5–25) after cancer diagnosis—all the patients except 1 (patient 2) had a metastatic disease, which included brain involvement in 2 cases (patients 1 and 3). ICIs used comprised nivolumab (3 cases), pembrolizumab (2 cases), and a combination of nivolumab and ipilimumab in 1 case. Median delay between ICI introduction and onset of the neurologic syndrome was 4 months (range: 2–8). When the neurologic syndrome ensued, the 2 patients with cerebral metastasis had stable or improved lesions on brain MRI, whereas the others showed no evidence of cancer dissemination in the CNS. Clinical syndromes included a combination of limbic encephalitis and diencephalitis (patients 1, 2, and 5), isolated limbic encephalitis (patients 3 and 6), and a syndrome characterized by ophthalmoplegia and motor neuron involvement (head drop) in patient 4. Onset of the neurologic symptoms was usually subacute (3 patients), whereas patient 5 had an acute onset and patient 1 a chronic/progressive course. All patients fulfilled the criteria for definite PNS.15 No statistically significant clinicodemographic differences (including sex, age at onset, cancer type, and main neurologic syndrome) were observed between our 6 patients and the overall cohort of 44 patients with anti-Ma2 Abs. During the study period 2015–2016, when the use of ICIs in France was at its starting point, we observed only 8 Ma2-PNS cases, meaning that a 112.5% increase was observed since ICI implementation in our center. Clinical and paraclinical results, together with associated complications in 3 patients, whereas in 1 case, it was attributed to the tumor progression. Of note, before ICI withdrawal, all patients except 1 (patient 4) showed a good response of cancer to immunotherapy, with stabilization or reduction of the neoplastic lesions. Intriguingly, for 1 case (patient 2), a serum sample was taken before ICI initiation and then stored in a biobank. The retrospective analysis of this sample revealed the presence of Ma2 Abs even before cancer immunotherapy onset. No antecedent serum sample was available for the other patients.

Impact of ICI use on the frequency of Ma2-PNS

We questioned whether ICI use had an impact on the frequency of Ma2-PNS in our center. During the study period 2017–2018, a total of 17 patients with Ma2 Abs were diagnosed in our reference center. The 6 Ma2-PNS described herein representing 35% of all the cases. During the biennium 2015–2016, when the use of ICIs in France was at its starting point, we observed only 8 Ma2-PNS cases, meaning that a 112.5% increase was observed since ICI implementation in France. As a matter of fact, the annual number of anti-Ma2 positivities diagnosed in our national reference center was relatively stable over the last decade, with a median of 4 cases per year (range: 1–6 cases), and this observation is therefore unprecedented. No other onconeural Ab targeting intracellular antigens demonstrated a similar increment (figure 2). A lower increment was demonstrated for the recently implemented neural surface Abs (such as NMDAR, Lgi1, CASPR2, GABAAR, and AMPAR), ranging from 30 to 50%, probably reflecting their relatively novel adoption in clinical practice if compared with the former group. Remarkably, only 1 case associated with neural surface antibody (CASPR2 positivity) developed after ICI initiation, accounting for 3% of all CASPR2 patients diagnosed in our center in the same period.

Discussion

We describe here 6 patients who developed anti-Ma2-PNS after receiving ICI treatment. Their demographic, clinical, and sequences. CSF analysis revealed inflammatory alterations in all cases, with the most common abnormality being an increased protein content (5 cases), followed by pleocytosis (2 cases) and presence of CSF-exclusive oligoclonal bands in 1 patient. Patient 5 showed additional anti-Ma1 positivity; no other onconeural Abs were detected in the remainder of the patients. The neurologic syndrome was moderately severe with a median pretreatment mRS score of 4 (range: 0–6), characteristic of a patient unable to walk unassisted and to attend to own bodily needs. Treatment included ICI withdrawal and corticosteroids in all patients. Additional treatment was adopted in 4 patients: 2 received IV immunoglobulin, 1 was treated with plasmapheresis, and 1 with rituximab. Median follow-up was 4 months (range: 1–6 months). Despite all these measures, 4/6 patients died, and the remaining 2 showed a moderate to severe disability (mRS score: 3 and 4, respectively). The cause of death was directly related to the neurologic involvement or its associated complications in 3 patients, whereas in 1 case, it was attributed to the tumor progression. Of note, before ICI withdrawal, all patients except 1 (patient 4) showed a good response of cancer to immunotherapy, with stabilization or reduction of the neoplastic lesions. Intriguingly, for 1 case (patient 2), a serum sample was taken before ICI initiation and then stored in a biobank. The retrospective analysis of this sample revealed the presence of Ma2 Abs even before cancer immunotherapy onset. No antecedent serum sample was available for the other patients.
<table>
<thead>
<tr>
<th>Patient no.</th>
<th>Sex, age (y)</th>
<th>Cancer (interval from diagnosis to ICI initiation)</th>
<th>ICI treatment (interval from ICI initiation to neurologic syndrome onset)</th>
<th>Neurologic symptoms/signs</th>
<th>Brain MRI, inflammatory alterations/type</th>
<th>CSF (white cells per mm³/ protein [g/L]/ OCB)</th>
<th>Ma2 Ab CSF (titer)/ serum (titer)</th>
<th>PNS treatment</th>
<th>mRS score before and after PNS treatment (length of follow-up)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>M, 79</td>
<td>Stage IV lung cancer (9 mo)</td>
<td>Pembrolizumab (2 mo)</td>
<td>Absent</td>
<td></td>
<td>n (0)/↑/NA</td>
<td>+ (1: 1,280)+ (1: 64,000)</td>
<td>Corticosteroids</td>
<td>5 → 3 (6 mo)</td>
</tr>
<tr>
<td>2</td>
<td>M, 71</td>
<td>Pleural mesothelioma (0.5 mo)</td>
<td>Nivolumab + ipilimumab (5 mo)</td>
<td>Narcolepsy-cataplexy, hyperphagia and weight gain (+12 kg over 6 mo), psychiatric symptoms</td>
<td>FLAIR hypersignal involving the uncus bilaterally, periventricular regions of the third ventricle and hypothalamus</td>
<td>↑/↑ (0.74)/−</td>
<td>+ (1: 1,280)+ (1: 1,024,000)</td>
<td>Corticosteroids Rituximab</td>
<td>3 → 6 (6 mo)</td>
</tr>
<tr>
<td>3</td>
<td>F, 57</td>
<td>Stage IV lung cancer (8 mo)</td>
<td>Nivolumab (8 mo)</td>
<td>Memory deficits, new-onset epilepsy, and psychomotor retardation</td>
<td>FLAIR bilateral MTL hypersignal</td>
<td>↑ (10)/↑ (0.79)/−</td>
<td>+ (1: 40,960)+ (1: 2,048,000)</td>
<td>Corticosteroids IVIG</td>
<td>3 → 6 (4 mo)</td>
</tr>
<tr>
<td>4</td>
<td>M, 47</td>
<td>Stage IV lung cancer (1 mo)</td>
<td>Pembrolizumab (8 mo)</td>
<td>Abrupt onset of diplopia Plateau: ophthalmoplegia + head drop</td>
<td>FLAIR bilateral MTL hypersignal</td>
<td>n (2)/n (0.38)/+</td>
<td>+ (NA)+ (NA)</td>
<td>Corticosteroids</td>
<td>4 → 6 (4 mo)</td>
</tr>
<tr>
<td>5</td>
<td>M, 55</td>
<td>Stage IV kidney cancer (25 mo)</td>
<td>Nivolumab (3 mo)</td>
<td>Abrupt onset of right ear hearing loss, followed by ataxia, vertigo, and memory deficits Plateau: hyperphagia, weight gain (+18 kg in 3 mo) excessive daytime sleepiness</td>
<td>FLAIR bilateral MTL hypersignal</td>
<td>n/↑ (0.92)/NA</td>
<td>+ (NA)+ (NA)</td>
<td>Corticosteroids PLEX</td>
<td>4 → 4 (3 mo)</td>
</tr>
<tr>
<td>6</td>
<td>M, 69</td>
<td>Stage IV lung cancer (5 mo)</td>
<td>Nivolumab (3 mo)</td>
<td>Confusion and focal seizures Plateau: confusion and decreased consciousness (GCS 13)</td>
<td>Absent</td>
<td>n (5)/↑ (0.60)/ NA</td>
<td>+ (1: 40,960)/ NA</td>
<td>Corticosteroids</td>
<td>4 → 6 (1 mo)</td>
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</table>

Abbreviations: FLAIR = fluid-attenuated inversion recovery; ICH = immune checkpoint inhibitor; IVIG = IV immunoglobulin; mRS = modified Rankin Scale; MTL = mesial temporal lobe; n = normal; NA = not available/not performed; OCB = oligoclonal band; PLEX = plasmapheresis.
paraclinical features were remarkably uniform. Most of them were male, middle-age/elderly adults who developed a neurologic syndrome characterized by prominent limbic and diencephalic involvement, mainly associated with lung cancer. This clinical presentation is in line with both the original descriptions of the Ma2 syndrome\(^9,10,16,17\) and the clinical features from the 44 remaining patients of our overall cohort. We, however, note 3 notable differences: (1) in the classic, paraneoplastic form, the neurologic syndrome is known to precede cancer diagnosis by several weeks to months. When the tumor is eventually found, it is usually detected at a limited disease stage. On the contrary, when the disease appeared as a complication of ICIs, patients already presented with metastasis and the neurologic syndrome manifested several months after cancer discovery. (2) Testicular cancer is a frequently associated neoplasia in Ma2-PNS, and patients with Ma2 autoimmunity in the context of testicular tumors tend to be younger. Because the current treatment of testicular cancer does not include ICIs, this could explain the older age at onset and higher percentage of lung cancer association observed herein. (3) Contrast enhancement of inflammatory brain alterations is usually detected in up to one-third of Ma2-PNS cases. However, this pattern was not observed in the cases elicited by ICI treatment.\(^6\)

Despite these differences, the inflammatory alterations detected by CSF analysis, the presence of well-characterized Abs, and the selective brain MRI involvement of the mesial temporal lobe and diencephalon structures strongly suggest

Figure 1 Results of paraclinical studies in patients with anti-Ma2 encephalitis triggered by immune checkpoint inhibitors [ICIs]

A. Limbic involvement  
B. Diencephalic involvement

Brain MRI (fluid-attenuated inversion recovery sequences) in 2 patients with anti-Ma2 encephalitis triggered by ICIs. Note the prominent limbic (A, axial view) and diencephalic (B, sagittal view) involvement (arrowheads).

Figure 2 Proportion of variation in antibody detection between 2017 and 2018 vs 2015 and 2016 at the French Reference Center for Paraneoplastic Neurological Syndromes

Note the 112% increase in Ma2-associated paraneoplastic neurologic syndrome detection observed after immune checkpoint inhibitor introduction. No other onconeural antibody (Ab) targeting intracellular antigens demonstrated a similar increment. A lower increment is observed for the recently implemented neural surface Abs, ranging from 30% to 50%, probably reflecting their relatively novel adoption in clinical practice if compared with the former group.
an immune-mediated pathogenesis. We therefore consider that the ICI treatment elicited the autoimmune encephalitis in our patients.

Because the anti–Ma2-associated syndrome is characterized by atypical manifestations such as increased daytime sleepiness, hyperphagia, and weight gain, it is important for the clinician to recognize the prominent features of this disease to avoid diagnostic pitfalls. These symptoms are related to the diencephalic involvement and need to be promptly differentiated from the clinical correlate of primary hypothyroidism, which is a much more common irAE that shares a similar presentation.22 As such, the use of the adoption of second-line immunosuppressants is probable in this situation, is not sufficient for all patients; (3) the adoption of second-line immunosuppressants is probably warranted for refractory cases.22 As such, the use of drugs—such as natalizumab—that can act on brain inflammatory processes without hampering the immune reaction against systemic localizations of cancer, was recently suggested.23

The pathogenesis of neurologic irAEs due to ICI use remains to be elucidated, although several lines of evidence suggest that (1) ICIs act by blocking the signaling from certain molecules—CTLA-4, PD-1, and its ligands—that exert inhibitory regulatory effects on T-cell activation, thus promoting antitumor immunity; (2) the antitumor immune response might in turn cross-react with CNS autoantigens leading to a PNS, as demonstrated in a preclinical model using CTLA-4 blockade; and (3) CD8+ T cells, activated by ICIs, were found to play a major effector role in neuronal death in PNS. In agreement with this model, it has been previously shown that the pathologic substrate of post-ICI encephalitis is characterized by prominent CD8+ lymphocytic perivascular infiltration.25

Two previous cases of ICI-induced anti-Ma2 encephalitis have been described. One concerned a patient with pleural mesothelioma treated with the anti-CTLA-4 Ab tremelimumab, and the second was in a patient with kidney cancer treated with nivolumab.25 Including the present series, this brings the total of cases reported in the literature up to 8. The reason for the increase in susceptibility to anti-Ma2 autoimmune response among all other Ab-associated PNS remains unclear. We propose that it reflects the fact that non–small-cell lung cancer is one of the cancers in which ICIs are most extensively used, and this tumor is known to associate with Ma2-PNS.26,27 We therefore hypothesize that an analogous increment of anti-Yo and anti-Ri syndrome will be seen after the adoption of ICIs in breast cancer.24,27 The same will probably occur for anti-Hu or anti-CV2/CRMP5 PNS if their use is extended to small-cell lung cancer.28,29

The retrospective detection of Ma2 Ab in the serum of one of our patients taken before ICI administration is an intriguing finding that deserves further discussion. First, it should be considered that at the time, the sample was taken and stored in a biobank, and the patient was asymptomatic. Neurologic symptoms appeared only 5 months later, following treatment with the combination of ipilimumab-nivolumab. Second, the finding of a confirmed Ma2 positivity in a patient without neurologic syndrome is exceedingly rare, whereas other onconeural Abs, such as anti-Hu and anti-CV2/CRMP5, are detected in 16% and 9% of neurologically asymptomatic patients with SCLC, respectively.15 These Abs are known to be reliable biomarkers of an underlying cancer but are not pathogenic because a T cell–mediated response is advocated as the cause of the neurologic syndrome.15,30 We therefore hypothesize, similarly to what we have observed in patients with ovary cancer with Yo-Abs and paraneoplastic cerebellar ataxia,31 that the tumor (a pleural mesothelioma in this case) expressed aberrantly the Ma2 antigens and triggered the systemic Ab production. This event per se was not sufficient to elicit a PNS, but required a loss of self-tolerance as permitted by the use of ICIs. This hypothesis needs to be verified in prospective studies assessing the presence and titer of onconeural Abs over time, their relation to immunotherapy, and the development of an overt neurologic syndrome. Our practical recommendation is to test patients undergoing ICI treatment for onconeural Abs before initiation of immunotherapy, and to closely follow those with an Ab positivity, with special caution for Ma2-positive cases. Patients with preexisting Abs are probably at an increased risk of developing irAEs, as demonstrated for anti-acetylcholine receptor autoantibodies and subsequent myositis in patients treated with avelumab.32

The present study is limited by its retrospective nature, small sample size, and, possibly, referral bias toward more complex and/or treatment-refractory cases. Nevertheless, it represents the 2-year experience of a national reference center focused on the diagnosis and treatment of PNS.
Discussion

We showed a clear association between ICI use in France and an increased frequency of anti-Ma2-PNS. Although final arguments proving a causal relationship between ICI and PNS development are lacking, there are several findings suggesting that this syndrome is related to ICI. Middle-aged/elderly men with lung cancer appeared to be at particular risk of developing post-ICI anti-Ma2-PNS. Given the anticipated rise in the use of immunotherapy for oncologic practices, we highlight the importance of early detection of these immune-mediated neurotoxic effects, which can be severe or even fatal.

Acknowledgment

The authors thank NeuroBioTec Hospices Civils de Lyon BRC (France, AC-2013-1867, NFS96-900) for banking sera and CSF samples. They also thank Dr. Pauline De L’Estang Du Rusquec, Dr. Youcef Douadi, Dr. Aurore Jourdain, Dr. Justin Le Tallec, and Dr Jerome Meunier who provided additional clinical data for the study. They gratefully acknowledge Véronique Rusquec, Dr. Youcef Douadi, Dr. Aurore Jourdain, Dr. Justin Le Tallec, and Dr Jerome Meunier who provided additional clinical data for the study. They gratefully acknowledge Véronique Rusquec, Dr. Youcef Douadi, Dr. Aurore Jourdain, Dr. Justin Le Tallec, and Dr Jerome Meunier who provided additional clinical data for the study.

Study funding

This study is supported by research grants from Agence Nationale de la Recherche (ANR-14-CE15-0001-MECANO), Fondation pour la recherche médicale (DQ20170336751).

Disclosure

The authors have no conflicts of interest to disclose. Go to Neurology.org/NN for full disclosures.

Publication history

Received by Neurology: Neuroimmunology & Neuroinflammation April 24, 2019. Accepted in final form July 8, 2019.

Appendix (continued)

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Increased frequency of anti-Ma2 encephalitis associated with immune checkpoint inhibitors
Alberto Vogrig, Marine Fouret, Bastien Joubert, et al.
Neurol Neuroimmunol Neuroinflamm 2019;6;
DOI 10.1212/NXI.0000000000000604

This information is current as of August 7, 2019