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Immune-Mediated Cerebellar Ataxias: Clinical Diagnosis and Treatment Based on Immunological and Physiological Mechanisms

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ABSTRACT

Since the first description of immune-mediated cerebellar ataxias (IMCAs) by Charcot in 1868, several milestones have been reached in our understanding of this group of neurological disorders. IMCAs have diverse etiologies, such as gluten ataxia, postinfectious cerebellitis, paraneoplastic cerebellar degeneration, opsoclonus myoclonus syndrome, anti-GAD ataxia, and primary autoimmune cerebellar ataxia. The cerebellum, a vulnerable autoimmune target of the nervous system, has remarkable capacities (collectively known as the cerebellar reserve, closely linked to plasticity) to compensate and restore function following various pathological insults. Therefore, good prognosis is expected when immune-mediated therapeutic interventions are delivered during early stages when the cerebellar reserve can be preserved. However, some types of IMCAs show poor responses to immunotherapies, even if such therapies are introduced at an early stage. Thus, further research is needed to enhance our understanding of the autoimmune mechanisms underlying IMCAs, as such research could potentially lead to the development of more effective immunotherapies. We underscore the need to pursue the identification of robust biomarkers.

Key Words Anti-GAD ataxia; Gluten ataxia; Immune-mediated cerebellar ataxias; Opsoclonus myoclonus syndrome; Paraneoplastic cerebellar degeneration; Postinfectious cerebellitis.

BACKGROUND: WHERE DO WE STAND?

History of immune ataxias: milestones

Immune-mediated pathophysiological mechanisms frequently affect the cerebellum, leading to progressive ataxia characterized by dysmetria in both motor and cognitive domains.1,2 The first documentation of patients with immune-mediated cerebellar ataxias (IMCAs) originated from Charcot.3 In a well-documented lecture on multiple sclerosis (MS) delivered in 1868, he described the presence of cerebellar ataxia (CA) in patients with MS, now known as the Charcot’s triad (intention tremor, scanning speech, and nystagmus). Another historical milestone was the first report of paraneoplastic cerebellar degeneration (PCD) by Brouwer4 in 1919, in which he described the association of CA with ovarian cancer.

In the 1980s, the identification of autoantibodies targeting cerebellar neurons led to a breakthrough in IMCAs. First, the discovery of anti-Yo antibody (Ab), an autoantibody described in a patient with ovarian cancer, demonstrated the immune nature of the insult resulting in PCD.5 This discovery was followed by the identification of specific autoantibodies, including anti-Hu, anti-Tr, anti-CV2, anti-Ri, anti-Ma2, and anti-VGCC Abs, often...
associated with specific types of neoplasms, including breast, uterine, ovarian, small cell lung carcinoma, and Hodgkin’s lymphoma. Second, the association of otherwise idiopathic CAs with autoantibodies against the cerebellum was also reported in patients without evidence of cancer. Two main clinical entities have been established since then, based on specific clinical features and types of associated autoantibodies: gluten ataxia (GA) by Hadjivassiliou et al. and anti-glutamic acid decarboxylase 65 Abs (GAD 65 Abs)-associated cerebellar ataxia (anti-GAD ataxia) by Honnorat et al.

Finally, in 2008, Hadjivassiliou et al. proposed a new clinical entity: primary autoimmune cerebellar ataxia (PACA). This entity encompasses all ataxic patients who exhibit features of IMCAs but with a serological profile that does not match any of the known main etiologies. Diagnostic criteria have been proposed recently. Thus, it has become easier to apply this diagnosis for such atypical patients to consider the use of immunotherapy.

**Classification**

IMCAs gather diverse etiologies, and the pathophysiology is not restricted to the cerebellum, also targeting cerebellar-related structures/connections in the central nervous system (CNS). The diversity renders the classification of IMCAs challenging. We previously proposed a classification based on 2 criteria: 1) whether the cerebellum is the main target of autoimmunity, and 2) whether autoimmunity is generated by a known trigger or not (Table 1). In MS, for example, the cerebellum is one of many targets within the CNS. Pure CA types, in which the cerebellum or its related structures are the sole target of autoimmunity, include GA, postinfectious cerebellitis (PIC), PCD, opsoconulus myoclonus syndrome (OMS), anti-GAD ataxia, and PACA. Thus, the term IMCAs is reserved for mainly pure or primarily CA, thus excluding more widespread autoimmune damage such as that seen in the context of MS. Although autoimmune limbic encephalitis can sometimes be characterized by CA in addition to cognitive/behavioral deficits and seizures, these conditions should not be categorized as IMCAs.

Pure or primarily CA etiologies are divided into two groups: 1) etiologies in which autoimmunity is triggered by other conditions, such as infection (e.g., PIC), neoplasm (e.g., PCDs), and gluten sensitivity (GA), and 2) etiologies in which autoimmunity is not triggered by any other condition (e.g., anti-GAD ataxia). When autoimmunity is triggered by other conditions, removal of the trigger should be the first line of therapy. This approach is important because categorization based on the autoimmune triggering factors can also offer therapeutic strategies.

In the case of PACA, immune-mediated mechanisms are strongly suspected, but the serological profile of the patients does not match any of the known diseases in IMCAs. Thus, this group is likely to be heterogeneous and probably includes several pathophysiological immune-mediated mechanisms.

**Prognosis and pathophysiology**

In keeping with the diverse etiologies, IMCAs show characteristically diverse clinical courses (Figure 1). These diverse clinical courses result in two problems when considering therapeutic strategies: the need for early intervention and the unclear immune-mediated pathophysiology.

**Cerebellar reserve and early intervention**

GA, PIC and the subacute type of anti-GAD ataxia show good response to immunotherapy (full or partial recovery and stabilization) (Figure 1). The good prognosis is in part due to the benign transient autoimmune insult, which is also seen in other autoimmune diseases. Alternatively, the reversibility can also be attributed to specific cerebellar capacity for compensation and restoration following the immune insult. We termed this capacity the cerebellar reserve. Various forms of synaptic plasticity and redundant mossy fiber-mediated inputs constitute the cerebellar reserve. Notably, the extent of progression of the pathophysiological mechanisms determines the degree of reversibility, suggesting the existence of a threshold. Although various immunotherapies can arrest the progression of the damage under this threshold, such intervention may not be accompanied by clinical improvement. Thus, immunotherapies should be introduced during the period when the cerebellar reserve is above this threshold. Such a threshold can be understood as the limit required to obtain a sufficient level of activity in the cerebellar circuitry.

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**Table 1. Classification of IMCAs**

| Cerebellar autoimmunity triggered by another disease or condition | (gluten sensitivity) |
| Cerebellar autoimmunity triggered by another disease or condition | (infection) |
| Miller Fisher syndrome | (infection) |
| Oposconulus myoclonus syndrome | (neoplasm) |
| Paraneoplastic cerebellar degenerations | (neoplasm) |
| Cerebellar autoimmunity not triggered by another disease or condition | *Anti-GAD ataxia* |
| Primary autoimmune cerebellar ataxia | Others |

*when cerebellar ataxias are the sole or main symptoms, the cerebellum is presumed to be the main target of autoimmunity. The term IMCAs is reserved for mainly pure or primarily cerebellar ataxia, thus excluding more widespread autoimmune damage, *opsoconulus myoclonus syndrome also includes idiopathic type, *anti-GAD ataxia occurs in other autoimmune backgrounds, such as neoplasm. IMCAs: immune-mediated cerebellar ataxias. GAD: glutamic acid decarboxylase. Adapted from Mitoma et al. Cerebellum 2016;15:213-232, under the terms of the Creative Commons CC BY license.*
Autoimmune mechanisms and response to treatment

The chronic subtypes of anti-GAD ataxia and PCD show poor prognosis (relapse and gradual or rapid progression in Figure 1), which reflects poor responsiveness to immunotherapy. The development of various autoimmune diseases can be attributed to common elementary processes, including autoimmunity triggered by deficits in immune tolerance or molecular mimicry, increased permeability of the blood-brain barrier or blood-nerve barrier, pathogenic actions of T cells (Th1/17 cells and CD8 T cells) or autoantibodies, and exacerbation of local neural inflammation (Figure 2). However, unlike the advances in the field of MS, little is known about therapeutic autoimmune targets in IMCAs.

In conclusion, the clinical entity of IMCAs has been established during the past three decades. To avoid missing therapeutic opportunities at the early stage, we should carefully promote IMCAs in the differential diagnosis of all ataxias. We also need more research directed toward the identification of new autoimmune pathogenesis-based therapies.

GENERAL ISSUES ON DIAGNOSIS AND IMMUNOTHERAPIES

Prevalence

Hadjivassiliou et al. (2017) investigated the prevalence of IMCAs on 1,500 UK patients with progressive ataxia. The authors reported that 30% of the patients had familial/genetic ataxia, although some did not show evident family history, and 9% of the patients had a cerebellar variant of multiple systemic atrophy (prevalence out of total progressive ataxic cases). Apart from the above, 25% had definite IMCAs; 20% had GA, and 2% had PCD, while 2% had anti-GAD ataxia, 1% had PIC, and <1% had OMS. Interestingly, 19% of the patients were classified as idiopathic sporadic ataxia. This category probably included a large number of patients with PACA. Among other groups, Gebus et al. (2017) reported a smaller number of IMCAs patients (2 with PCD and 1 patient with PIC among 80 patients). In a systematic study of 684 South Korean patients with progressive ataxia, only 21 had IMCAs; 14 patients with PCD, 3 patients with PIC, and 4 patients other immune

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**Figure 1.** Clinical courses of immune-mediated cerebellar ataxias. (A) Six patterns of clinical courses of immune-mediated cerebellar ataxias. (B) Schematic diagram of the concept of the cerebellar reserve. PIC: postinfectious cerebellitis, GA: gluten ataxia, OMS: opsoclonus myoclonus syndrome, Anti-GAD: anti-glutamic acid decarboxylase ataxia, PCD: paraneoplastic cerebellar degeneration. Adapted from Mitoma et al. Curr Neuropharmacol 2019;17:33-58, with permission of Bentham Science Publishers.
causes. The discrepancy might be attributed to a number of different factors: The high prevalence of IMCAs in Hadjivassiliou’s study was due to a large number of patients with GA. GA was first characterized and reported by this group and there is therefore the potential for referral bias. Another important factor is the serological test utilized to make a diagnosis of GA (please see section on GA). More recently, the prevalence of GA in Asian countries has been found to be higher than was previously estimated. For example, patients with GA have recently been reported in Japan. A survey in China also showed that the prevalence rates of positive anti-transglutaminase 6 (TG6) IgA were 35% in 100 sporadic CA patients and 9.8% in 51 healthy controls (p = 0.001). Another factor is the so-called “idiopathic” unknown etiology diagnosis. As discussed above, we have argued that this group includes a large number of patients with PACA. The newly established diagnostic criteria on PACA will be helpful in resolving this issue.

Further studies are needed to examine the prevalence of IMCAs or GA in progressive CAs. However, early diagnosis and intervention is critical during a period when the cerebellar reserve is preserved. In this regard, clinicians should always consider the possibility of IMCAs in the differential diagnosis of patients developing CAs.

Clue to diagnosis

Patients with IMCAs exhibit common clinical manifestations in spite of divergent etiology. The onset of CA is usually acute/subacute or sometimes chronic/insidious. The presence of other autoimmune disorders is common in some patients. The main clinical feature is gait ataxia, which influences standing and steady walking and is associated with mild limb incoordination and dysmetria, oculomotor ataxia, and scanning speech. In keeping with these symptoms, MRI characteristically shows vermian atrophy, which can be a clue for the differential diagnosis from degenerative or genetic CAs. As the pathology progresses, atrophy becomes more evident, preferentially affecting the vermis. CSF studies sometimes show the presence of pleocytosis or oligoclonal bands. There may be a context of systemic impaired immunity (affecting skin, joints, or other organs).

Autoantibodies

As discussed earlier in the section on the history of IMCAs, identification of autoantibodies is important in the diagnosis (Table 2). Immunohistochemistry may reveal the binding of autoantibodies toward cerebellar components; therefore, if available, it can add further evidence of the diagnosis. Autoantibodies in IMCAs can be divided into two categories: 1) autoantibodies...
suggestive of specific etiologies, and 2) nonspecific autoantibodies found in other neurological and systemic conditions, including CAs, which provide only possible autoimmune pathophysiology. The former types of autoantibodies include antigliadin and TG6 Abs in GA, and anti-Yo, Hu, CV2, Ri, and Ma2 Abs for PCD.

The nonspecific autoantibodies are also subdivided into two subcategories: autoantibodies that are assumed to have pathogenic roles and the nonpathogenic autoantibodies (i.e., a diagnostic marker).

The majority of the pathogenic autoantibodies include autoantibodies toward ion channels or ion channel-related proteins (Ca²⁺ or K⁺ channel), glutamate receptors, or GABA synthesis enzyme (GAD), some of which have been confirmed as pathogenic in both in vitro and in vivo preparations. Impairments of neural excitability and synaptic transmissions can occur throughout the nervous system, such as limbic encephalitis. We previously proposed a nomenclature based on the pathogenic role of the autoantibody, i.e., the pathogenic Ab to be coined to the name of the clinical entity (e.g., anti-GAD ataxia).

Additionally, some autoantibodies are reported to be associated with CAs in only a few patients. These autoantibodies are classified into a category of not-well-characterized autoantibodies. The significance of these autoantibodies remains to be determined, but their presence can be a clue to the autoimmune etiology of the ataxia.

**Principles of treatment**

When autoimmunity is caused by another trigger, priority should be given to the treatment of the underlying condition: for example, gluten-free diet (GFD) in GA and surgical excision or chemotherapy and radiotherapy of the neoplasm in PCD (Figure 4). For other IMCAs, immediate immunotherapy is recommended (Figure 4).

Depending on the rapidity of progression, induction immunotherapy can be used to arrest progression of CA, followed by maintenance immunotherapy to prevent relapse. Various immunotherapies have been used, ranging from intravenous immunoglobulins (IVIg), glucocorticosteroids, mycophenolate, plasmapheresis, and rituximab, either alone or in combination, and the selection is often based on the etiology and presentation. These therapies should be introduced during the period when the cerebellar reserve is still preserved.

To establish a therapeutic rationale, the following two issues need further consideration. First, to date, there are no large-scale randomized studies comparing therapeutic strategies for each IMCA. Second, the effects of each therapeutic intervention should be quantified with more sensitive biomarkers if possible at an early stage of the disease when MRI shows no atrophy. Recent studies highlighted the potential usefulness of MR spectroscopy as a sensitive biological marker of the response to treatment in CA patients. The ratio of N-acetylaspartate (NAA)/creatine (Cr) area is decreased in patients with CAs (Figure 3B) relative to the control and increases in those who respond to immunotherapy and exhibit clinical improvement.

**PARTICULAR ISSUES IN COMMON ETIOLOGIES**

This section focuses on main etiologies in IMCAs (Table 3).

**Gluten ataxia**

GA is defined as sporadic CA associated with gluten sensitivity. The autoimmunity is triggered by the sensitivity to gluten found in wheat, rye and barley. Gluten, a family of protein containing grains, is composed of gliadin and glutenin.

**Diagnosis and treatment**

GA affects individuals in their 40–50s and exhibits either chronic or insidious onset. Gluten-sensitive enteropathy (celiac disease) is seen in approximately half of the patients. The human leukocyte antigen (HLA) type DQ2 is detected in 70% of the patients, and association with other autoimmune diseases, such as thyroiditis, type 1 diabetes mellitus, and pernicious anemia, is common. Patients present with gait ataxia and variable degree of limb ataxia, scanning speech and, consistently, MRI shows vermian involvement. GA can be associated with CSP abnormalities, with oligoclonal bands in up to 50% of patients. The association of sensorimotor axonal neuropathy and focal myoclonus, rarely palatal tremor, is sometimes observed.
Diagnosis is based on the presence of serological evidence of gluten sensitivity, which is assessed using autoantibody testing. Native anti-gliadin Ab (AGA) remains to be the most reliable test for the diagnosis. However, the cutoff level for the diagnosis of GA should be noted. In GA patients without enteropathy, the main autoimmune response occurs within the CNS, not the periphery, resulting in low levels of serum AGA. The calibration process used in certain commercially available AGA kits is based on the use of serum from patients with celiac disease (CD) and, therefore, the cutoff level is too high for the detection of GA. Anti-transglutaminase 2 (TG2) Ab, found in patients with CD, is negative in 53% of GA patients free of enteropathy.
An expert dietitian should be considered before switching to immunotherapy. However, anti-TG6 Ab is positive in 72% of GA patients on the basis of AGA. Since TG6 is primarily expressed in the CNS, anti-TG6 Ab could be an important and possibly more specific biomarker.

GFD is the first line of therapy for GA since it can eliminate antigens that can trigger immune-mediated mechanisms, similar to the strategy used in CD. One large-scale study based on 43 patients with GA showed significant improvements in CA and a decrease in AGA in patients who adhered to GFD, compared with those who did not. Other reports described the effectiveness of IVIg in nonresponders to GFD. However, it is now considered that the lack of response to GFD is due to either poor adherence to GFD or hypersensitivity to gluten, where a small amount of gluten present in commercially available gluten-free food or due to cross-contamination can cause strong autoimmune reactions perpetuating the cerebellar damage. Persistently high levels of AGA are present in the above two groups of patients. On MR spectroscopy, patients on strict GFD show an increase in the ratio of NAA/Cr area in the cerebellar vermis, whereas no such increase is seen in patients on GFD with persistently positive antibodies and those who do not adhere to GFD. Taken together, in patients with persistently high AGA titers or those who do not adhere to GFD, maintenance therapy with IVIg or immunosuppressants (e.g., rituximab) is recommended.

Autoimmune pathophysiological mechanisms

Loss of immune tolerance is considered the main mechanism, as in coeliac disease. Digested gliadin is deamidated and cross-linked by TG2, leading to the creation of an immunostimulatory epitope for HLA-DQ2 or HLA-DQ8 on antigen-presenting cells. These epitopes are presented to CD4+ T cells, from which cytokines are released to facilitate the production of antibodies against gliadin and TG2 (TG2-gliadin complex). The neural damage appears to be due to, at least in part, the actions of autoantibodies against tissue transglutaminase, a calcium-dependent enzyme located in intra- and extracellular spaces. Both anti-TG2 and anti-TG6 Abs recognize and cross-react with neurons. Intraventricular administration of anti-TG2 or anti-TG6 Abs elicits ataxia in mice. TG2 activity is upregulated through calcium binding, and anti-TG2 Abs bind preferentially to the calcium-activated enzyme conformation. Taken together, antibody-induced deficits of TG2 in its enzymatically active conformation can cause cellular dysfunction. The role of TG6 in cerebellar function is also supported by the observation that mutations in the TGM6 gene can cause CA (spinocerebellar ataxia type 35).

Although the mechanisms of cell-mediated brain damage remain uncertain, the above immunological studies add support to the therapeutic rationale of gluten avoidance in food and reductions of anti-TG2/6 and AGAs.

Myoclonic ataxia with refractory celiac disease

CA associated with cortical myoclonus is a rare subtype of gluten sensitivity-related neurological disorder. Although the myoclonus is of cortical origin, hyperexcitability of the cerebral cortex is elicited by cerebellar dysfunction. Characteristically, this subtype shows resistance to GFD. The neurological refractoriness is associated with residual enteropathy, which is detected by repeat duodenal biopsies. All such patients require GFD plus immunosuppression, usually with mycophenolate and in some instances cladribine, in addition to anti-epileptic drugs for the control of the myoclonus. The prognosis remains poor.

Postinfectious cerebellitis

PIC is defined as cerebellar inflammation induced by immune-mediated mechanisms triggered by viral or bacterial infection. Some groups have termed this type of autoimmune ataxia as para-infectious cerebellitis or acute cerebellar ataxia. However, when the cerebellar inflammation is caused by direct invasion of viral or bacterial microorganisms, it is called acute cerebellitis.

Diagnosis and treatment

PIC affects mostly young children after an episode of infection, usually viral infection, most commonly varicella. Other viruses, such as Epstein-Barr virus, Coxsackie virus, influenza A and B virus, parainfluenza virus, rubeola virus (measles), ru-
| Table 3. Clinical profiles of common etiologies of immune-mediated cerebellar ataxias |
|------------------|------------------|------------------|------------------|------------------|------------------|------------------|
|                  | Gluten ataxia    | Postinfectious   | Opsoclonus myoclonus | Paraneoplastic   | Anti-GAD ataxia  | Primary autoimmune |
|                  |                  | cerebellitis     | ataxia syndrome     | cerebellar       |                  | cerebellar ataxia |
| Prevalence among all progressive cerebellar ataxias\(^{18}\) | 20%              | 1%               | 0.8%              | 3%               | 2%               | Unknown (among 20% of idiopathic sporadic ataxias) |
| Autoimmune background |                 |                  |                  |                  |                  |                  |
| Trigger of autoimmunity | Gluten ingestion | Postinfectious, children: varicella. Adults: EBV, mycoplasma | Paraneoplastic (neuroblastoma) postinfectious, primary autoimmune | Neoplasm (breast, uterus, ovarian, small cell lung carcinoma, Hodgkin’s disease and others) | Unknown | Unknown |
| HLA | Type DQ2 or DQ8 | - | - | - | - | Type DQ2 |
| Well or partially characterized antibodies | Anti-gliadin (IgG/IgA), anti-TG2, TG6 | None | Anti-Ri (for paraneoplastic) | Anti-Yo (53%), anti-Hu (15%), anti-CV2 (4%), anti-Ri (2%), anti-MA2 (2%) anti-Tr (5%)\(^{9}\) | Anti-GAD65 (high titer) | None |
| Less well characterized autoantibodies | - | Anti-Gluδ2 | Anti-glycin R (for paraneoplastic) | Anti-VGCC (2%), anti-SOX1, anti-ZIC4, PCA-2, anti-Homer3, anti-CARP VIII, anti-PKCy, anti-Ca/ARHGAP26, anti-mGlenuR | - | Anti-Cerebellum (immunohistochemistry) anti-GAD65 (low titer), anti-Homer3, anti-Gluδ2 |
| Associated autoimmune diseases | Celiac disease (47%), thyroid, type 1 diabetes mellitus, pernicious anemia | - | - | - | Type 1 diabetes, thyroid, pernicious anemia | Thyroid, Sjögren’s, type 1 diabetes, primary biliary cirrhosis, pernicious anemia, vitiligo |
| Clinical profile |                 |                  |                  |                  |                  |                  |
| Time course | Insidious and chronic | Acute | Subacute | Subacute | Insidious and chronic or subacute | Insidious and chronic |
| Age and gender | 40–50s, females (55%) | Mainly in children, rarely adults | Mainly in children, rarely adults | 50–60s, | 60s, females (mostly) | 50s |
| Main manifestations of cerebellar involvement | Gait ataxia | Gait ataxia | Opsoclonus, myoclonus in addition to gait ataxia | Pancerebellar ataxia | Gait ataxia | Gait ataxia |
| Associated neurological symptoms | Cortical myoclonus in some, neuropathy | - | - | - | Epilepsy, ophthalmoplegia and Stiff-Person syndrome | - |
| CSF abnormalities | Oligoclonal bands in some cases | High WBC count. High IgG levels in 50% of patients. Oligoclonal bands in some | Sometimes; high WBC count and protein level | Frequently; elevation in WBCs and protein, oligoclonal bands | Sometimes; CSF oligoclonal bands | Not studied |
| Cerebellar atrophy on MRI | Present depending on duration of ataxia before diagnosis | None | None | Not at onset but may develop rapidly | Present depending on duration of ataxia | Present depending on duration of ataxia |

EBV: Epstein-Barr virus, HLA: human leukocyte antigen, WBC: white blood cell, TG: transglutaminase, VGCC: voltage gated calcium channel, CASPR2: contactin-associated protein-like 2, mGlueR: metabotropic glutamate receptor, GAD65: glutamic acid decarboxylase 65, ZIC4: zinc finger protein of the cerebellum 4, PCA2: purkinje cell antibody 2, GluR52: glutamate receptor delta2, PKCy: protein kinase C gamma, Ca/ARHGAP26: Ca/Rho GTPase-activating protein 26, SOX1: sex determining region Y-related high-mobility group box 1, CARP VIII: carbonic anhydrase-related protein VIII. Adapted from Mitoma et al. MedLink Neurology 2019 Aug 23 (www.medlink.com), with permission of MedLink.\(^{11}\)
Bulavirus (mumps), and rubella virus (German measles), and bacteria, such as Corynebacterium diphtheriae (diphtheria), Bordetella pertussis (whooping cough), Salmonella typhi (typhoid fever), Legionella (Legionnaires disease), Leptospira (leptospirosis), and Mycoplasma (mycoplasmosis) can be involved. \(^7,^8,^10,^50,^52\) Borrelia (Lyme disease) is also implicated. One large-scale study based on 73 patients \(^53\) showed that 60% of the patients were between 2 and 4 years of age, and 25% of these patients had varicella, 52% had other viral infections, and 3% developed PIC after immunization. The mean latency between infection and the onset of CAs was 9.9 ± 7.9 days. However, some patients (19%) showed no definite preceding infection.

Patients develop mainly acute-onset afebrile gait ataxia, meningeal signs, high intracranial pressure with or without extracerebellar manifestations, such as temporary clouding of consciousness, seizures, altered mental status (e.g., extreme irritability), or extracerebellar focal signs. \(^53\) The presence of these clinical features is often suggestive of direct infective etiology rather than immune-mediated mechanisms (PIC). \(^32\) Additionally, mild behavioral changes, such as mild irritability, hyperactivity, moodiness and whining, are sometimes noted by parents and often correlate with the severity of CA. \(^52\) These manifestations could reflect cerebellar cognitive and emotional disorders. CSF studies show pleocytosis in most of the patients (roughly equal numbers of granulocytes and lymphocytes in approximately 75% of the patients, and sometimes lymphocyte predominance in the remaining cases) and high CSF/serum IgG index in half of the children. Oligoclonal bands are sometimes present. \(^53\) MRI is usually negative, with no signs of atrophy or abnormal intensity areas. \(^53\) Nevertheless, adult patients can develop cerebellar atrophy after several months.

Since PIC is self-limiting, close observation of the patient is recommended. \(^10,^11\) One large-scale study reported full recovery of CAs in 72% of 60 pediatric patients within two months. \(^53\) Only when the CAs persist or progress are various combinations of immunotherapeutic agents available for use. \(^10,^11,^50\)

Autoimmune pathophysiological mechanisms

Autoimmune mechanisms remain obscure. For example, it is not certain whether molecular mimicry, observed in Guillain-Barré syndrome (GBS, a typical infection-induced neuroimmune disease), also operates in PIC. Although the lack of cerebellar atrophy and self-limiting clinical course suggests immune-mediated cerebellar functional disorders, the exact mechanisms remain unclear. A recent study \(^26\) described the unique actions of anti-GluRδ Ab, which is associated with PIC in some patients. \(^26\) In one experimental study, injection of an antibody toward the putative ligand-binding site of GluRδ2 resulted in endocytosis of AMPA receptors and attenuated synaptic transmission and ultimately led to the development of ataxic behavior in mice. \(^54\) Interestingly, the endocytosis of AMPA receptors, which has a transient nature, could explain the reversibility in PIC. Thus, fur-
ther studies are needed to understand how autoimmune reactions transiently disturb cerebellar function.

**Opsoclonus myoclonus syndrome**

OMS has divergent categories, including paraneoplastic, postinfectious, and idiopathic. It affects mainly children and is especially associated with neuroblastoma. Infiltration is characteristically evident in paraneoplastic OMS, while cell loss and extensive T cell infiltration appears to occur not only within the cerebellum but also in inhibitory controls from the cerebellar cortex on cerebellar nuclei neurons. It is assumed that the net inhibitory output from Purkinje cells (PCs) in the cerebellar cortex is attenuated in both opsinoclonus and action myoclonus. In ocular motor controls, this elicits hyperactivation of the fastigial nucleus neurons, which in turn facilitates the activities of excitatory burst neurons, leading to a chaotic cascade of events. In the cerebro-cerebellar loop, the pathway of the dentato-thalamo-cortical tract is hyperactivated, in which even subthreshold inputs can easily activate the motor cortex, generating the action myoclonus in limb movements. Consistently, action myoclonus is of cortical origin, and a low correlation between the cerebellum and motor cortex is observed in functional MRI.

In conclusion, there is no explanation on how the autoimmune processes, typically triggered by diverse pathologies, such as neoplasm and infection, commonly produce a unique functional deficit, decrease the inhibitory output from the cerebellar cortex and generate hyperexcitability of their targets.

**Paraneoplastic cerebellar degeneration**

PCD comprises cerebellar degeneration caused by neoplasm-triggered autoimmunity. The presence of characteristic onconeural autoantibodies is not only diagnostic for PCD but also provides a clue to the specific type of neoplasm underlying PCD (Table 2). Although aggressive chemotherapies and immunotherapies have been attempted, the prognosis remains poor unless the underlying cancer can be completely and rapidly eradicated.

**Diagnosis and treatment**

PCD shows acute or subacute onset and is sometimes preceded by prodromal clinical symptoms such as nausea, vomiting, and dizziness, resembling viral infection-related disease. Subsequently, patients show gait ataxia, which is followed by pancerebellar involvement. CSF studies show changes consistent with inflammation, including moderate lymphocytic pleocytosis, high protein concentrations, high IgG index, and oligoclonal bands. Identification of the characteristic autoantibodies helps in the establishment of diagnosis, although seronegative cases have been described. Since subacute CAs are listed as classical symptoms defined in the 2004 diagnostic criteria, a definite diagnosis is based on 1) confirmation of the presence of cancer, which develops within 5 years of the diagnosis of CA, or 2) appearance of well-characterized onconeural Abs (anti-Yo, anti-Hu, anti-CV2, anti-Ri, anti-MA2). CA is the first manifestation of neoplasm in up to 70% of the patients with PCD, highlighting the need for systemic evaluation for malignant tumors following the identification of onconeural autoantibodies.

Following confirmation of the diagnosis, anti-neoplasm ther-
apies should be introduced immediately to prevent metastasis and remove antigen(s) driving the autoimmune response. The selection of cancer treatment, such as surgery, radiotherapy and/or chemotherapy, is based on the type of the neoplasm. Regardless of the selected cancer treatment modality, various types of immunotherapies (single or combinations of corticosteroids, IVlg, plasmapheresis, immunosuppressants, and rituximab) should be administered. In spite of these treatment protocols, the prognosis is poor, although therapeutic benefits have been reported in some case reports, and better prognosis was noted in patients with anti-Tr Ab. The study of Peterson et al. on 22 patients with anti-Yo Ab reported therapeutic improvement in less than 10% of the trials (one in four trials of plasmapheresis, one in 17 trials of corticosteroids, and none in four trials of cyclophosphamide). Furthermore, Rojas et al. also reported no benefits from the combinations of immunotherapies in their study of 23 patients with PCDs.

There have been three long-term studies on survival time. Candler et al. examined the clinical courses of 63 patients with paraneoplastic neurological syndromes, including 13 patients with PCDs. Three of the 13 (23%) patients with PCDs died before the final follow-up, and the mean survival time from onset of neurological symptoms was 42 months (95% confidence interval: 32–52). Keime-Guibert et al. reported a long-term follow-up study of 16 patients with PCDs. When the clinical benefits were determined as a more than one point change in the Rankin Score, none of their patients met this criteria (three patients showed stabilization). The median survival time from the first therapy was 10.2 months (range 2–38). Fourteen of the 16 (88%) patients died during the follow-up. The cause of death was cancer-related in 9 patients, neurological-related in 3 patients, and unknown in the remaining patients. Shams’ili et al. examined the relationship between effectiveness of treatment and associated antibodies in their study of 50 patients with PCDs. The median survival times from diagnosis were 113 months in patients positive for anti-Tr Ab, 69 months in those positive for anti-Ri Ab, 13 months in those positive for anti-Yo Ab, and 7 months in those positive for anti-Hu Ab. It should be noted that even patients with anti-Tr Ab suffered severe disability or dependence. Taken together, the survival time is generally short once the diagnosis is established, excluding a few patients with anti-Tr Ab.

The poor prognosis could be attributed to the remnant tumor tissue and metastasis, which could cause persistent activation of the immune system and strong autoimmune response. Currently, clinical trials aimed at stopping the progressive autoimmune insults are being conducted. In one such trial, the combination of IVlg or plasmapheresis with cyclophosphamide was reported to be effective in a subgroup of patients.

**Autoimmune pathophysiological mechanisms**

Pathological studies have identified various inflammatory changes in the early stages of PCD, including perivascular cuffing by lymphocytes, infiltration of lymphocytes within the PC layer, and microglial activation. The infiltrates consist of both T and B cells, plasma cells, microglia/macrophage lineages, and reactive Bergmann cell gliosis. A similar infiltration, albeit of lower magnitude, was found in the dentate nucleus and the pons. Disease progression is associated with massive loss of PC without inflammation (inflammatory cells or deposits of immunoglobulins and complements). Since the loss of PC is often very rapid, such a burnout stage can be observed in some patients at an earlier stage.

Yo (Cdr2) and Hu onconeural antigens are intracellularly located in tumor and neuronal cells. This favors molecular mimicry mechanisms. First, whether autoantibodies that target intracellular antigens Cdr and Hu proteins play pathogenic roles in the development of the inflammatory cerebellar lesions is still a matter of debate today. Previous studies showed that anti-Yo Ab toward Cdr2 was internalized and induced cell death through apoptosis. In this regard, Cdr2 encodes a leucine zipper motif and interacts with another leucine zipper motif on c-Myc, the nuclear transcription factor, to regulate nuclear levels of c-Myc and c-Myc-associated signaling activities. Cdr2-specific anti-Yo Ab inhibits the interaction of Cdr2 with c-Myc, possibly allowing excess c-Myc to enter the nucleus and impair cell cycle signaling involved in cell apoptosis. The binding of anti-Yo Ab to Cdr2 can also disrupt interaction with the mortality factor-like proteins MRGX and/or NFkB, leading to changes in transcriptional activity. Another study showed that administration of anti-Hu and Yo induced the expression of adhesion molecules and more intense cell differentiation rather than cell death. Furthermore, another histochemical study showed a lack of both IgG deposits and B cell infiltration. A study of passive transfer of onconeural Abs that targeted intracellular antigens and immunization using protein or DNA failed to induce ataxic symptoms in animals. Thus, many researchers believe that antibody-mediated mechanisms are not the main autoimmune mechanisms underlying PCDs associated with anti-Yo and anti-Hu Abs.

Patients with anti-Yo (Cdr2) Ab or anti-Hu Ab show accumulation of Cdr2- or Hu-specific T cells in blood and CSF, respectively. Notably, the lymphocytes found in tumors, peripheral blood, and affected nervous tissue are thought to originate from the same clone population. The infiltrated lymphocytes in the cerebellum are mainly CD3+ and CD8+ T cells and only a few are CD4+ lymphocytes. In general, CD4+ and CD8+ T cell interaction is necessary to induce inflammation, where CD4+ T cells exhibit a Th1 phenotype and secrete IFN-γ and TNF-α. These studies suggest that PCDs are mediated mainly...
by a CD8+ T-cell immune response toward an autoantigen recognized by onconeural Ab. The cytotoxic lymphocytes appear to be aggressive toward cells presenting Cdr2 or Hu antigens and possibly induce cell death through one or more of the following mechanisms; 1) activation of TNF and FasL receptors, which triggers the apoptotic cell death cascade, and 2) excretion of granules that destroy cells through the actions of granzyme-B, perforin, and T-cell restricted intracellular antigen-1 (Figure 6).

Despite the above evidence suggesting the involvement of cell-mediated mechanisms in the development of PCDs, the transfer of lymphocytes from patients with anti-Yo Ab did not elicit any PC damage. In addition, although all small cell lung carcinoma expresses the Hu antigen, and a clinically useful tumor immune response can be detected in up to 20% of the patients, the paraneoplastic neurologic syndrome is extremely rare, suggesting the robust tolerance to neural onconeural antigens in vivo.

Based on this notion, Blachère et al. argued that CD8+ T cells that target intracellular antigens, in conjunction with CD4+ T cells, can induce tumor autoimmunity, but not neurologic autoimmunity. They proposed that 1) CD8+ T cells can only activate the peripheral immune response and 2) CD8+ T cells can break CNS tolerance to induce PCDs if they interact with both CD4+ T cells and autoantibodies. This interaction between cell- and autoantibody-mediated mechanisms might provide the rationale for the design of new therapies that can target T and B cells using tacrolimus and rituximab, respectively.

The following mechanisms can also underlie the break of CNS tolerance: 1) structural modification of certain intracellular proteins, which stimulates the immune system, 2) death of cancer cells and release of intracellular antigens with subsequent exposure to the immune system, 3) persistence of the above autoimmune responses to neoplasm causes secretion of cytokines that elicit vascular hyperpermeability and infiltration of immune cells, and 4) simultaneous downregulation of regulatory T cells.

**Anti-GAD ataxia**

GAD is an enzyme that catalyzes the conversion of glutamate to GABA. GAD has two isoforms: GAD65 and GAD67. The autoantibodies against the smaller isozyme are associated with CA. Serum and CSF titers of anti-GAD65 Ab are higher in anti-GAD ataxia, at usually more than 10,000 U/mL (or 10 to 100-fold higher), compared to those of patients with type 1 diabetes mellitus (T1DM). The triggering factor of this autoimmunity is not clear at present. Anti-GAD ataxia can be associated with other types of IMCAs, such as PCD and GA. While autoimmunity toward GAD65 affects the entire CNS, the cerebellum is one of the most vulnerable areas.

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**Figure 6.** Possible autoimmune mechanisms underlying paraneoplastic cerebellar degenerations. Naive CD4+ T cells are first sensitized by onconeural antigens (for example, Cdr2 or Hu) in the peripheral lymph nodes, and they differentiate into antigen-specific Th1 or ThFH cells. Activated CD8+ T cells are aggressive toward cells presenting these onconeural antigens through FasL receptor-mediated or perforin/granzyme-mediated apoptosis. It is uncertain how CNS tolerance is broken down to develop cerebellar degenerations.
Diagnosis and treatment

The condition mostly affects women in their 60s, with subacute or chronic/insidious time course. Some patients show evidence of other autoimmune conditions, such as T1DM, autoimmune thyroid diseases, and pernicious anemia. Anti-GAD ataxia is sometimes associated with extracerebellar symptoms, including epilepsy, ophthalmoplegia and Stiff-Person syndrome. The clinical course can be either subacute or chronic. CSF studies sometimes show oligoclonal bands, whereas MRI shows normal or atrophic changes depending on the duration of illness. The diagnosis is confirmed by the presence of high titers of intrathecally produced anti-GAD Ab.

The serum level of anti-GAD antibodies in the context of the diagnosis of anti-GAD ataxia merits further consideration. There is currently no consensus on the threshold used for “low titer” and “high titer” levels of anti-GAD antibodies. In cases of anti-GAD Ab more than 2,000 U/mL (high titer), one can safely consider anti-GAD ataxia. However, titers in the range of 100–200 U/mL in an appropriate clinical/biological/neuroimaging context may be observed in immune ataxias. Some patients with CA and anti-GAD Ab less than 100 U/mL have been reported to show benefits to immunotherapies. Anti-GAD Ab less than 100 U/mL may raise a suspicion of autoimmune diathesis and require therapy. It is also important to keep in mind that various laboratories use different techniques for the assays.

The aim of any induction therapy should be to minimize progression of CA. Thus, various immunotherapies, alone or in combination (more common), are used until remission. Both induction and maintenance therapies include corticosteroids, IVIg, immunosuppressants, plasmapheresis, and rituximab, either alone or in combination. There appear to be no significant differences in the therapeutic benefits among the above immunotherapies. The prognosis is better in the subacute type than in the chronic type of anti-GAD ataxia.

Autoimmune pathophysiological mechanisms

The significance of anti-GAD65 has been a matter of debate. Some researchers have argued that anti-GAD65 Ab have no pathogenic roles in the development of CAs based on the following reasons: 1) these actions were abolished after the absorption of anti-GAD65 Ab using recombinant GAD65 and anti-GAD65 Ab elicited no actions in slices from GAD65 knockout mice where inhibitory transmissions were mediated compensatorily by GAD67, 2) the b78 monoclonal Ab with epitope-specificity in CAs has pathogenic actions, whereas another monoclonal Ab has no such actions in T1DM, and 3) differences in neurological phenotype might also be related to epitope specificity. Notably, low titer anti-GAD Ab has no pathogenic actions.

Internalization and dissociation of GAD65 with vesicles

Anti-GAD65 Ab is internalized, presumably during exocytosis or endocytosis. Since exposure of GAD65 is assumed to be during exocytosis, anti-GAD65 could access GAD65, although the exact mechanisms are still not clear. Anti-GAD65 Ab is reported to impair the association of GAD65 with vesicles, resulting in deficits in GABA packaging into vesicles and shunting of vesicles to the release sites (Figure 7).

From functional disorders to cell death

A decrease in GABA release attenuates the spill-over GABA-induced presynaptic inhibition on glutamate release from neighboring parallel fibers, resulting in a major imbalance between GABA and glutamate and excitotoxicity (Figure 8). Consistently, one autopsy study showed the complete loss of PCs.

Clinical correlation

Previous studies showed that some patients respond well to immunotherapies and that the clinical improvement in CA correlates well with the fall in Ab titers, suggesting that Ab titers better reflect functional disorders rather than cell death. The peak CA might be influenced by secondary cell-mediated mechanisms.

Taken together, the above experimental studies clearly show that anti-GAD65 Ab could serve as a trigger of CA (Figure 7). These studies also provide a generalized principle regarding an-
tibody-induced neurological diseases; autoantibodies are polyclonal, and a cocktail of these polyclonal autoantibodies determines the clinical profile, depending on the proportions of pathogenic epitopes. In addition, pathomechanisms underlying the excitotoxicity could be generalized in other pathologies, such as ischemic diseases and degenerative CAs (Figure 8).

Primary autoimmune cerebellar ataxia

Although the clinical profile spectrum of PACA meets the features of IMCAs (e.g., subacute onset, autoimmune disease history, predominant gait ataxia, association of autoantibodies, and good benefits from immunotherapies), a subgroup of patients present with features that do not meet the criteria of the above common subtypes.\(^9,10,11,14,15,17,18\) Thus, the spectrum of PACA serves as an umbrella that covers heterogeneous etiologies.\(^9,10,11,14,15,17,18\) Consistently, various types of less characterized autoantibodies are associated in some patients (Table 2), and immunohistochemical studies show a variety of staining patterns (60% of the patients).\(^14\) Despite the heterogeneity, patients with PACA sometimes show HLA type DQ2, which is observed in certain autoimmune diseases (CD, GA, T1DM, Stiff-Person syndrome, autoimmune thyroid disease and autoimmune polyendocrine syndromes), but not in healthy subjects.\(^14,15\)

Many types of immunotherapies have been used in PACA. A small number of studies have reported the effectiveness of IVlg, prednisolone, plasmapheresis or rituximab in 9/19 patients at the early stage of PACA.\(^119\) Progression to wheel-chair dependence was faster in patients with neuronal nuclear and/or cytoplasmic antibody than those positive for plasma membrane protein antibody,\(^114\) suggesting the need for the selection of therapeutic strategy based on the underlying autoimmune pathoetiiology. In this regard, MR spectroscopy of the cerebellum could be an important tool in monitoring the response to immunotherapies.\(^15,117,118\)

**OTHER RARE ETIOLOGIES IN IMCAs**

This section focus on rare etiologies in IMCAs.\(^117,118\)

**Miller Fisher syndrome**

In 1956, Fisher\(^120\) proposed a new clinical entity of an acute-onset triad of ophthalmoplegija, ataxia and areflexia syndrome, which is currently termed Miller Fisher syndrome (MFS). MFS is assumed to be a representative disease of infection-triggered autoimmunity, similar to GBS.\(^10\) A large-scale study involving 50 consecutive patients showed that viral infection (usually respiratory infection) or bacterial infection (usually *Campylobacter jejuni*) preceded the appearance of the triad manifestations, with a median interval of 8 days.\(^121\) High serum titers of the specific marker anti-GQ1b Ab are found in up to 90% of patients with MFS, but not in those with other autoimmune neuropathies, such as GBS and chronic inflammatory demyelinating polynemopathy. Anti-GQ1b Ab cross-reacts with epitopes present in the liposaccharide of MFS-associated *Campylobacter jejuni* strains, suggesting molecular mimicry.\(^112\)

MFS shows self-limiting clinical course, thus requiring no immunotherapy.\(^10\) One large-scale study showed full recovery within 6 months in all patients.\(^121\) The speed of recovery and the final outcome were the same in patients who received corticosteroids, IVlg, or plasmapheresis and those who were treated conservatively.\(^121\) The good prognosis suggests that any autoimmune-mediated impairment in MFS is reversible. It remains to be determined whether the autoimmune response functionally attacks peripheral or central structures. Based on the findings of GQ1b expression in large-diameter dorsal ganglion neurons\(^12\) and the presence of enhanced lesions in the spino-cerebellar tracts at the lower medulla on MRI,\(^123\) several researchers have argued for the presence of peripheral lesions in MFS. Moreover, cerebellar involvement has been confirmed more directly in a number of recent studies. One study using FDG-PET showed hypermetabolism in the cerebellum and brainstem,\(^124\) and another study...
using MR spectroscopy showed abnormal signals in the cerebellum during the course of the illness. Furthermore, immunocytochemical studies using sera of MFS patients confirmed the presence of autoantibodies to antigens in the molecular layer. Recently established clinical entities

Dipeptidyl-peptidase-like protein-6 (DPPX), an auxiliary subunit of the Kv4.2 potassium channel, and contactin-associated protein-like 2 (Caspr2), a protein associated with the Kv1 potassium channel, can be targets of autoimmunity. Since potassium channels are distributed throughout the CNS, CAs are one of the diverse features of these etiologies. Patients with anti-DPPX Ab show multifocal neurological symptoms such as agitation, mild confusion, hyperekplexia, myoclonus, trunk stiffness and CAs, which are attributed to the hyperexcitability induced by potassium channel deficits. Recently, CAs with myoclonus were also reported as the sole manifestation of this entity. Additionally, encephalitis associated with anti-Caspr2 Ab is defined as ≥ 3 core manifestations, including encephalic signs, CAs, peripheral nerve hyperexcitability, dysautonomia, neuropathic pain, insomnia, and weight loss. Notably, stereotypical episodes of paroxysmal CAs (mimicking episodic ataxias) are found in some patients with anti-Caspr2 Ab. The gait imbalance, limb ataxia, and slurred speech usually last a few minutes to a few days and usually improve following immunotherapy.

Figure 8. Overview of the possible synaptic consequences of decreased GABA release in the cerebellum. (Normal GABA regulation) In normal conditions, GABA released from GABAergic neurons binds to GABA<sub>A</sub> receptors located on PC and induces an IPSC. GABA also diffuses out of the synapse and binds to GABA<sub>B</sub> receptors, reducing the release of the excitatory neurotransmitter glutamate. Glutamate ultimately activates NMDA receptors on excitatory neurons and is regulated in part through NO. Glutamate is taken up by astrocytes via the EAAT pathway. (Functional impairment) When GAD65 Ab is produced intrathecally, GAD65 Ab may bind GAD65 and interfere with GABA release. Reduced GABA levels increase glutamate levels as a consequence of the lower inhibition of GABAB receptors. Continuously high glutamate levels can develop divergent effects, as detailed in the following. Glutamate activates microglia, which in turn releases more glutamate. Saturation or impairment of EAAT will reduce glutamate re-uptake by astrocytes. Activation of x<sup>c</sup>- increases the extracellular glutamate release and finally, the NMDA/NO feedback regulation fails. All these effects result in an increase in glutamate concentrations. (Cerebellar atrophy) The excessive Ca<sup>2+</sup> influx stimulates calpain I and nNOS, leading to mitochondrial dysfunction, ER stress and DNA damage. In this scheme, Purkinje cells and granule cells are assumed. However, NMDA receptors are expressed in granule cells, whereas metabotropic glutamate receptors are expressed in Purkinje cells. Mossy fibers strongly excite granule cells via NMDA and AMPA receptors, contributing to cerebellar neuronal hyperactivity. GA: GABA<sub>A</sub> receptors, GB: GABA<sub>B</sub> receptors, GAD: glutamic acid decarboxylase, A: AMPA receptors, N: NMDA receptors, M: metabotropic glutamate receptors, EAAT: excitatory amino acid transporters, Blue dots: GABA, Red dots: glutamate, Glu: glutamate, Gln: glutamine, GS: glutamine synthetase, (-): inhibitory effects.
These two novel entities suggest that the cerebellum is one of the vulnerable areas in patients with potassium channel dysfunction. Myelin-associated glycoprotein (MAG) plays an important role in the maintenance of the myelin sheath, not only in Schwann cells but also in the CNS. The association of anti-MAG Ab with distal demyelinating neuropathy is well established in patients with monoclonal gammopathy of unknown significance. Recently, CA associated with anti-MAG Ab was described in five patients. All patients had IgM gammopathy, and four of the five showed clinically evident neuropathy. MR spectroscopy showed cerebellar involvements, which diminished after improvement of CAs with rituximab.

In contrast to these autoantibody-related etiologies, chronic lymphocytic inflammation with pontine perivascular enhancement responsive to steroids (CLIPPERS) is characterized by marked perivascular T cell infiltration. The lymphocytes, mainly CD4-dominant lymphocytes, infiltrate the perivascular area and more diffusely in the white matter of the pons and adjacent rhombencephalic structures, such as the cerebellar peduncles, cerebellum, medulla, and the midbrain. The perivascular infiltration appears as characteristic gadolinium enhancement on MRI, with multiple “punctate” and/or “curvilinear” gadolinium-enhancing lesions, resulting in “peppering” of the pons with or without peripontine lesions. The conditions affect mainly men in their 50s. The patients show subacute onset of the combination of brainstem symptoms and CAs, which include pan cerebellar ataxias, dysarthria, dysphagia, dysgeusia, oculomotor abnormalities, altered facial sensation, facial nerve palsy, vertigo, pyramidal signs and sensory disorders. CLIPPERS is characterized by responsiveness to corticosteroids. The initial treatment is intravenous methyl prednisolone followed by maintenance immunotherapy using the combination of oral prednisolone and corticosteroid-sparing immunosuppressants. Since the clinical course seems to be relapsing–remitting in nature, long-term maintenance therapy is required.

CONCLUSION

IMCAs include diverse etiologies, suggesting that the cerebellum can be the target of many types of autoimmune responses with different pathophysiological mechanisms. The identification of new clinical entities (anti-DPPX ataxia, anti-Caspr2 ataxia, anti-MAG ataxia, and CLIPPERS) suggests that further novel entities are likely to be discovered in the near future. IMCAs of certain etiologies respond well to immunotherapy at least in the early stage, while the cerebellar reserve is preserved. Thus, clinicians should identify therapeutic opportunities early. Conversely, a subgroup of etiologies shows poor prognosis, thus requiring a future breakthrough in therapeutic strategies. Further advances in our understanding of these complex immune mechanisms are important in dictating future immunotherapies.

Conflicts of Interest

The authors have no financial conflicts of interest.

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Author Contributions

Conceptualization: Hiroshi Mitoma. Data curation: Hiroshi Mitoma. Formal analysis: Hiroshi Mitoma. Investigation: Hiroshi Mitoma. Project administration: Hiroshi Mitoma, Mario Manto. Resources: Marios Hadjivasiliiou. Supervision: Mario Manto, Marios Hadjivasiliiou. Validation: all authors. Writing—original draft: Hiroshi Mitoma. Writing—review & editing: Mario Manto, Marios Hadjivasiliiou.

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