

Topic: Autoimmune neurological diseases associated with autoantibodies specific for synaptic antigens

Encephalitis associated with autoantibodies binding to γ -aminobutyric acid-A, γ -aminobutyric acid-B and glycine receptors: immunopathogenic mechanisms and clinical characteristics

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ABSTRACT

Recent discoveries of neural antibodies have facilitated the diagnosis of immune-mediated, immunotherapy-responsive neurologic disorders. Antibodies that target inhibitory central nervous system receptors, such as γ -aminobutyric acid-B, γ -aminobutyric acid-A, and glycine receptors, disrupt inhibitory regulatory synaptic functions, and lead to neuronal hyperexcitability. The myriad of neurologic manifestations associated with these antibodies includes seizures, encephalopathy, muscle rigidity and stiffness. This article provides a review of the immunopathogenic mechanisms and the clinical and therapeutic implications of autoimmune encephalitis associated with these antibodies that target inhibitory receptors.

Key words: Autoimmune encephalitis; autoimmune epilepsy; limbic encephalitis; neural antibodies

INTRODUCTION

Recent discoveries of neural antibodies that bind to antigenic targets in the brain have led to a paradigm shift in the clinical approach to patients presenting with encephalopathy,^[1-4] cognitive change,^[5] and refractory seizures.^[4,6] With a wider availability of neural antibody testing, a significant proportion of the patients, who were previously diagnosed with encephalitis of undetermined etiology have been shown to have neurologic symptoms caused by an underlying autoimmune disorder and some of these patients respond favorably to immunosuppressive treatments.^[7] Neural antibodies that target channels or receptors on the neuronal cell surface can interfere with the function of these proteins, leading to altered neuronal excitability, and a myriad of neurologic syndromes that

mirror genetic and pharmacologically induced disorders of the target receptors.^[8] In this review, we describe the immunopathogenic mechanisms of autoimmune encephalitis associated with antibodies targeting the inhibitory synaptic receptors γ -aminobutyric acid-B (GABA_B), γ -aminobutyric acid-A (GABA_A), and glycine receptors (GlyRs), together with their clinical and therapeutic implications.

ANTI-GABA_B RECEPTOR AND ANTI-GABA_A RECEPTOR ENCEPHALITIS

GABA, the main inhibitory neurotransmitter in the brain, binds to metabotropic and ionotropic receptors to regulate neuronal activity. To date, GABA_B receptor (GABA_BR) and GABA_A receptor (GABA_AR) have been identified as antigenic targets of autoimmunity [Table 1].

Anti-GABA_BR encephalitis

The GABA_BR is a metabotropic G-protein-coupled

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receptor on presynaptic, postsynaptic, and extrasynaptic membranes, expressed in both the central and peripheral nervous systems, particularly the hippocampus, thalamus, and cerebellum.^[9] This receptor is a heterodimer comprising two subunits: GABA_{B1} and GABA_{B2}. Both subunits need to be co-expressed in order to form a functional receptor.^[9,10] The extracellular domain of the GABA_{B1} subunit binds to GABA while the GABA_{B2} subunit couples the receptor with the effector G protein.^[11] Antibodies to GABA_BR bind to the B1 subunit of the GABA_BR,^[4] the component that is required for GABA binding and receptor function.

GABA_BR exert inhibitory regulatory effects on synaptic transmission by inhibiting presynaptic voltage-gated calcium channel-mediated neurotransmitter release and by activating postsynaptic potassium channels, resulting in hyperpolarization of neuronal membranes, and inhibition of adenylate cyclase.^[11,12] GABA_BR dysfunction is implicated in a variety of neurological disorders such as epilepsy, in which genetic mutations may play a role.^[13,14] Pharmacological disruption of the GABA_BR leads to seizures, cognitive deficits, and behavioral changes,^[10,15,16] all of which may be seen in autoimmune anti-GABA_BR limbic encephalitis.

Antibodies to GABA_BR were first described in 15 patients with limbic encephalitis, in whom subacute early onset of seizures was a distinctive feature.^[4] Seizures were predominantly of temporal lobe onset with secondary generalization, and 3 of the 15 patients developed status epilepticus. Memory impairment, confusion, hallucinations, and behavioral changes consistent with limbic involvement were frequently seen. Electroencephalography (EEG) changes included epileptiform discharges, electrographic ictal activity, and/or temporal lobe slowing.

Magnetic resonance imaging (MRI) brain imaging typically demonstrates unilateral or bilateral increased T2/fluid attenuated inversion recovery signal changes in the medial temporal region, consistent with limbic encephalitis.^[4,17] Extratemporal changes in the grey and white matter, cerebellum, basal ganglia, and brainstem have also been reported.^[4,17-19] Cerebrospinal fluid (CSF) examination may yield lymphocytic pleocytosis, elevated protein, and oligoclonal bands.^[4] As with other types of limbic encephalitis, the EEG, MRI brain, and/or CSF exam may be normal, and should not preclude the diagnosis or presumptive treatment when the clinical presentation is suspicious.

Limbic encephalitis, the most common neurologic manifestation of anti-GABA_BR encephalitis,^[4,20,21] typically occurs in the setting of very high GABA_BR antibody titers.^[4,20] A widening phenotypic spectrum of anti-GABA_BR disorders including cerebellar ataxia,^[17,20,22,23]

opsoclonus-myoclonus,^[17,18] and brainstem encephalitis^[19] is now appreciated. Extralimbic presentations may be explained by the high expression of GABA_BR outside of the hippocampus, including the cerebellum.^[9] Other rarer neurological features of anti-GABA_BR encephalitis include chorea, myelopathy, peripheral neuropathy, and myopathy, particularly in patients with lower antibody titers.^[4,20] It is uncertain, whether all of these extralimbic neurologic manifestations can be attributed solely to the GABA_BR antibodies or whether the co-existence of other neural autoantibodies may contribute to the increasingly diverse neurological features being reported. Known neural antibody accompaniments to the GABA_BR antibodies include the 65 kDa isoform of glutamate decarboxylase (GAD-65), voltage-gated calcium channels (N-type and P/Q type), voltage-gated potassium-complex (VGKC-complex), and neuronal nuclear and cytoplasmic antibodies [such as antineuronal nuclear autoantibody (ANNA-1), ANNA-3, collapsing response mediator protein-5 IgG, and anti-glial nuclear autoantibody/SOX-1 antibodies].^[4,20-22]

In comparison to other cell surface neural antibody associated encephalitis, anti-GABA_BR encephalitis is probably uncommon. In a clinical service laboratory, only 7 of 3,989 (0.2%) patients with suspected autoimmune encephalopathy were found to have the GABA_BR antibody.^[20] There is no obvious gender predisposition for this neurologic disorder. A paraneoplastic etiology is diagnosed in approximately half of the patients with GABA_BR antibodies.^[4,17,20,21] The neurologic disorder usually precedes the diagnosis of malignancy, and the most frequently encountered tumor is small cell lung cancer.^[4,20] Tumors are more likely to be detected in older patients.^[4] Although, lung tumors from patients with GABA_BR encephalitis have not been studied for GABA_BR expression, samples of archived small cell lung cancers from patients without encephalitis were found to react with both guinea pig and human GABA_B IgG. This suggests that the GABA_BR could be expressed by small cell lung cancer and could potentially trigger an autoimmune reaction.^[4] Other oncologic associations of anti-GABA_BR encephalitis include neuroendocrine lung tumor, multiple myeloma, esophageal carcinoma, malignant melanoma, and carcinoid of the thymus.^[4,19,20,22,23]

Neurological improvement has been reported in up to 90% of patients with anti-GABA_BR encephalitis who received immunotherapy and appropriate cancer treatment (if the tumor was detected).^[4,17,20,21] As the cases reported so far were retrospectively ascertained, there was heterogeneity in the immunotherapies used. A variety of immunotherapies have been used successfully, including various combinations of first-line agents, corticosteroids, intravenous immunoglobulin (IVIg),

and/or plasma exchange (PLEX).^[4,20] Patients who do not respond to these treatments warrant second line and maintenance treatments, such as rituximab, cyclophosphamide, mycophenolate mofetil, and azathioprine.^[17,20,23,24] Neurological improvement may be incomplete or not sustained. Despite optimal immunosuppressive treatment, patients with GABA_BR antibodies can deteriorate due to tumor progression, chemotherapy-related complications, and/or treatment-resistant relapses.^[4,17,22-24] Further studies to elucidate the optimal treatment regimens are needed. The presence of an underlying small cell lung cancer and the co-existence of other paraneoplastic neural antibodies targeting intracellular (neuronal nuclear and cytoplasmic) antigens have been suggested as poor prognostic indicators.^[17,20]

Anti-GABA_AR encephalitis

The GABA_AR is a ligand-gated ion channel located at synaptic and extrasynaptic sites that functions to mediate fast inhibitory synaptic transmission.^[25,26] Activation of the GABA_AR triggers opening of intrinsic chloride channels, thereby eliciting an inhibitory postsynaptic potential.^[27] Disruption of GABA_AR results in increased neuronal excitability and seizures.^[27] Mutations in the $\alpha 1$ and $\beta 3$ subunits of the GABA_AR gene have been implicated in epilepsy syndromes.^[27-29] Benzodiazepine and barbiturate, medications used for the treatment of seizures and status epilepticus, enhance GABAergic inhibition to exert an anticonvulsant effect.^[30]

GABA_ARs are pentamers comprising combinations of five subunits that form chloride ion channels. Different combinations of subunits result in functional heterogeneity. Synaptic GABA_ARs, which contain the α ($\alpha 1-3$), β , and γ subunits, are responsible for phasic inhibition. By contrast, extrasynaptic and perisynaptic GABA_ARs, which are responsible for tonic inhibition, comprise $\alpha 4$ or $\alpha 6$ subunits combined with β and δ subunits.^[31] The GABA_AR antibody binds to the $\alpha 1, \beta 3$, or both subunits of the synaptic GABA_AR.^[32,33] GABA_AR antibodies reduce the density of the GABA_AR at synaptic sites when applied to rat hippocampal neurons, suggesting that antibody binding leads to the relocation of GABA_AR from the synaptic membrane.^[32,33] This phenomenon is similar to the loss of synaptic GABA_AR and resultant neuronal hyperexcitability observed in epilepsy and status epilepticus.^[27] The combined reinforcing effects of antibody-mediated synaptic GABA_AR relocation, together with the status epilepticus-induced loss of GABA_AR, could support a postulated model to explain the severity of seizures in patients with anti-GABA_AR encephalitis.^[33]

Recently, 18 patients with autoimmune encephalitis and prominent seizures were described with GABA_AR

antibodies, 6 of whom had very high antibody titers.^[33] Patients with high titers in both serum and CSF developed a particularly rapid, severe progressive encephalopathy with refractory seizures and/or status epilepticus, for which intensive care admission for pharmacologically-induced coma was required. Other reported clinical manifestations in GABA_AR antibody seropositive patients are opsoclonus-myoclonus, affective problems, hallucinations, mutism, aphasia, memory impairment, hemiparesis, chorea, cerebellar ataxia, and Stiff-man syndrome (SMS).^[32,33] GABA_BR, GAD-65, N-methyl-D-aspartate receptor (NMDAR), leucine-rich, glioma-inactivated 1 and contactin-associated protein-like 2 antibodies frequently co-exist in these patients.^[32,33] A propensity for other neurological autoimmune conditions such as myasthenia gravis has also been noted.^[32,33]

GABA_AR antibodies are reported in both children and adults (age 2-74 years), but larger cohorts need to be characterized.^[33] A low frequency of tumors in seropositive patients has been reported. In the initially published study of 18 patients, only one patient was found to have cancer (Hodgkin lymphoma).^[33] Two recent additionally reported cases had invasive thymoma.^[32]

The electroencephalograms of patients with anti-GABA_AR encephalitis may demonstrate generalized slowing suggestive of encephalopathy, multifocal ictal and interictal discharges, or status epilepticus.^[33] CSF findings range from normal to lymphocytic pleocytosis.^[33] Distinctive to GABA_AR antibodies, the majority of patients, especially those with high antibody titers, had extensive temporal and extratemporal MRI brain abnormalities^[33] which could be a consequence of autoimmune inflammation in the brain or prolonged ictal activity. The extensive radiologic changes contrast with those of patients with limbic encephalitis associated with other neuronal synaptic and cell surface antibodies, such as NMDAR and VGKC-complex antibodies, in which MRI abnormalities are often confined to the mesial temporal regions.

Despite the severity of their presentation, 80% of the patients reported with anti-GABA_AR encephalitis demonstrate partial or complete recovery with a combination of immunotherapy, antiepileptic drugs, and supportive treatment.^[32,33] In severe cases, multiple immunotherapies may be required. Treatment options are the same as with GABA_BR and GlyR antibody mediated disorders. In addition to immunotherapy, early recognition and treatment of epilepsy, as well as supportive treatment (including ventilation support) are pivotal.

GlyR antibody encephalitis

Glycine, a key neurotransmitter for fast postsynaptic

Table 1: Key demographic, clinical, and AI/oncologic associations of the GABA_BR, GABA_AR, and GlyR- α 1 antibodies

	GABA _B R	GABA _A R	GlyR- α 1
Clinical features	Limbic encephalitis (memory impairment, hallucination, confusion, behavior changes) with early and prominent seizures Less commonly: cerebellar ataxia, opsoclonus-myoclonus, brainstem encephalitis, chorea, myelopathy, peripheral neuropathy, and myopathy	High serum antibodies concentration: rapidly progressive encephalopathy, refractory seizures, and status epilepticus Low serum antibodies concentration: seizures, SMS, opsoclonus-myoclonus, behavioral change, psychosis, confusion, chorea, ataxia, hallucinations, and hemiparesis	Axial/limb spasms, rigidity, and myoclonus (SMS) + brainstem signs (PERM) Optic neuropathy, seizures, cognitive impairment, autonomic disturbance, respiratory failure, and transverse myelitis also reported in isolation or with SMS or PERM
Onset	Mostly subacute/acute	Acute/subacute	Subacute > acute > chronic
Age group	Wide range, children, and adults	Wide range, children, and adults	Wide range, children, and adults
Gender, male:female	1.3:1	2:1	1:1
Inflammatory CSF*	73%	40%	50%, but OCB frequently negative
MRI	65% abnormal (medial temporal > extratemporal changes)	High serum antibodies concentration: 100% multifocal temporal and extratemporal T2/FLAIR hyperintensities	< 30% T2/FLAIR abnormalities in temporal lobes. SC abnormalities rarely reported (short, multifocal, and LETM)
EMG	-	-	60% abnormal (continuous motor activity, stimulus induced motor activity)
Other coexisting neural antibodies	56% (VGCC, AGNA, GAD-65, VGKC-complex, NMDAR, ANNA-1, -2 and -3, CRMP-5 IgG, amphiphysin, BRSK2)	70% (AChR, NMDAR, GABA _B , GAD-65, VGKC-complex)	Rare (GAD-65, MOG, NMDAR, aquaporin-4, VGKC-complex)
Associated tumors	62% (SCLC most common, also neuroendocrine lung, malignant melanoma, esophageal, malignant melanoma, thymus anaplastic carcinoid)	15% (invasive thymoma, Hodgkin lymphoma)	Approximately 10% to date (breast cancer, lymphoma (both Hodgkin and NHL), leukemia, lung cancer, melanoma)
Fatalities (%)	36	20	< 10

*Inflammatory CSF includes pleocytosis, high protein, raised IgG index, and/or oligoclonal bands. AChR: acetylcholine receptor; AI: autoimmune; AGNA: anti-glia nuclear autoantibody; ANNA: antineuronal nuclear autoantibody; BRSK2: BR serine/threonine-protein kinase-2; CRMP-5: collapsing response mediator protein 5; CSF: cerebrospinal fluid; EEG: electroencephalogram; EMG: electromyography; FLAIR: fluid attenuated inversion recovery; GABA_AR: γ -aminobutyric acid receptor A subunit; GABA_BR: γ -aminobutyric acid receptor B subunit; GAD-65: glutamic acid decarboxylase-65; GlyR- α 1: glycine receptor alpha-1 subunit; LETM: longitudinally extensive transverse myelitis; MOG: myelin oligodendrocyte glycoprotein; MRI: magnetic resonance imaging; NHL: nonHodgkin lymphoma; NMDAR: N-methyl-D-aspartate receptor; OCB: oligoclonal bands; PERM: progressive encephalomyelitis with rigidity and myoclonus; SC: spinal cord; SMS: stiff-man syndrome; SCLC: small cell lung carcinoma; VGCC: voltage-gated calcium channel; VGKC-complex: voltage-gated potassium channel complex

inhibitory neurons in the CNS, has a complex functional pathway that involves pre- and post-synaptic GlyR interacting with other neurotransmitters (GABA and glutamate), NMDAR, and postsynaptic anchoring proteins like gephyrin. Antibodies directed at any of these targets may affect the glycinergic system, resulting in neurological dysfunction.^[34] Whether due to strychnine (a GlyR antagonist) poisoning, genetic mutations of the GlyR gene (hereditary hyperekplexia), or immune-mediated encephalitis, GlyR dysfunction may be associated with severe muscle spasms, stiffness, agitation, seizures, myoclonus, autonomic instability, and/or respiratory failure.^[35,36]

GlyRs, pentamers of α 1- α 4 and β -subunit proteins, are ligand-gated chloride ion channels, widely distributed in the CNS. They are predominantly expressed in the olfactory bulb, retina, hippocampus, brainstem (auditory, visual, vestibular, and sensory nuclei), cerebellum, and spinal cord.^[37,38] Glycine binding mediates opening of the GlyR chloride channel, resulting in hyperpolarization of the membrane potential and reduced neuronal excitability. The GlyR antibody targets the α 1 subunit of the postsynaptic GlyR and is associated with hyperexcitable neurologic disorders [Table 1].^[39] Gephyrin allows multiple GlyR to cluster together on

the synaptic membrane. Antibodies to gephyrin, an anchoring protein in the postsynaptic GlyR, have been described only in a single case to date.^[40]

Classic neurologic manifestations associated with GlyR antibodies are progressive encephalomyelitis with rigidity and myoclonus (PERM) and SMS.^[39,41,42] PERM and SMS were first described as separate clinical entities, but today these 2 conditions are considered to belong to a continuum of CNS hyperexcitability disorders. Patients with PERM and SMS share common features of rigidity, painful spasms, autonomic disturbances, hyperekplexia, and myoclonus. The widespread distribution of hyperexcitability and brainstem involvement classically distinguishes PERM from SMS and associated psychiatric symptoms such as anxiety are more commonly observed in SMS patients.^[43-46] The autoimmune nature of these conditions, and specifically the involvement of GlyR antibodies in some cases was not appreciated until recently.^[41,42] SMS was initially associated with antibodies to GAD-65 (60-70% of cases),^[47,48] gephyrin (1 case),^[40] and amphiphysin (< 5% cases, in the setting of both small cell lung and breast cancers).^[48,49] In 2008, Hutchinson *et al.*^[41] reported the first case of PERM with GlyR antibodies. It remains unresolved whether all of these

antibodies are truly pathogenic.^[50,51] McKeon *et al.*^[42] reported that 10 of 81 (12%) patients with SMS spectrum disorders were positive for GlyR antibodies. Interestingly, GlyR seropositivity was associated with better responsiveness to immunotherapy regardless of GAD-65 status, suggesting a pathogenic role. There are no passive transfer animal models of GlyR antibodies to date.^[39,42]

The spectrum of GlyR antibody encephalitis manifestations is now widening beyond classic PERM and SMS. Various combinations of psychiatric disturbances, cognitive dysfunction, seizures (focal/generalized epilepsy and new-onset status epilepticus), and movement disorders, autonomic instability with central hypoventilation, pseudobulbar and/or oculomotor dysfunction, steroid responsive optic neuropathy, and transverse myelopathy have now been described with GlyR antibodies.^[39,52-54]

The onset of symptoms in GlyR antibody neurologic syndromes is typically acute to subacute. In the largest case series of 52 GlyR antibody positive patients with a variety of presentations, there appears to be no sex predominance and all age groups are vulnerable.^[39] Patients frequently have a history of other autoimmune disorders.^[39] As with other autoimmune encephalopathies, GlyR antibodies may co-exist with other antibodies, such as NMDAR, GAD-65, VGKC-complex, myelin oligodendrocyte glycoprotein, and aquaporin-4 antibodies although this is rare.^[39,42,53,55,56] Tumors are identified in less than 20% of cases (thymoma, lymphoma, breast cancer, small cell lung carcinoma and leukemia).^[39,42,57,58]

GlyR antibodies may be detected in both serum and/or the CSF.^[39,59] CSF evaluation is possibly more sensitive than serum, therefore testing both is recommended.^[42] CSF lymphocytic pleocytosis or raised protein may be seen, and oligoclonal bands were negative in 50-70% of 2 case series recently published.^[39,56] Imaging is typically normal. Rarely, MRI temporal lobe T2-weighted abnormality with subsequent hippocampal volume loss is detected, particularly in cases associated with significant seizure activity. EEG may be normal, or show features of focal or generalized ictal activity.^[59]

A combination of immunotherapies (corticosteroids, IVIg, PLEX, cyclophosphamide), pharmacological therapies targeting symptoms of motor hyperexcitability and pain (clonazepam, diazepam, baclofen, gabapentin), and anticonvulsants (levetiracetam) are required to control clinical symptoms.^[39,42] Eighty percent of patients with GlyR antibodies showed a substantial response to immunotherapy.^[39,42] Two cases were reported that responded dramatically to thymectomy in addition to other immunotherapy.^[39,60] In the largest case series

to date, 6 out of 52 (12%) patients continued to have sporadic relapses whilst on treatment.^[39]

CONCLUSION

An increased awareness of the autoimmune mechanisms underlying cases of noninfective encephalitis and/or refractory seizures has led to increased recognition, earlier treatment, and improved outcomes in a subgroup of patients previously considered untreatable. Antibodies targeting the inhibitory receptors GABA_B, GABA_A, and glycine are three more recently appreciated, but important antibodies to consider in refractory seizure disorders and encephalitis of unclear etiology. A high index of suspicion and an awareness of the expanding clinical spectrum of these antibody-mediated disorders should prompt early neural antibody testing in patients with typical constellations of neurological symptoms, in particular refractory seizure disorders and encephalitis of unclear etiology. Once identified early, these conditions may be responsive to immunotherapy. There are sparse data to recommend one immunotherapeutic regime over another. Large cohort studies of patients with anti-NMDAR encephalitis suggest that first line therapy should comprise corticosteroids, IVIg, and/or PLEX, followed by second line immunotherapy (cyclophosphamide, rituximab, or both) in patients who fail to respond to initial treatment.^[61] A practical approach, guided by the literature on autoimmune encephalitis with antibodies against neuronal surface antigens, is suggested in Figure 1. Immunotherapy needs to be complemented by supportive, symptomatic medical therapy. There is a consensus that early treatment confers better outcomes. Age and antibody appropriate tumor screening should be performed in all cases and may be aided by testing for other co-existing neural antibodies.^[62]

The neurologic hyperexcitability effects of antibody binding to GABA_A, GABA_B, and GlyRs (and potentially other receptors in the future) reflect the important functions mediated by these inhibitory neuronal synaptic receptors. More research is needed in order to better understand this novel category of immune-mediated encephalitis. Further studies could focus on immunopathogenic mechanisms of these antibodies in causing disease, as these may be potential targets for directed treatment. To date, the numbers of patients reported with these antibodies remain small, with most cases retrospectively identified. With increasing access to testing for neural antibodies, the clinical spectrum of these autoimmune encephalitides may continue to expand. Systematic studies of prospectively identified, newly diagnosed cases should help to provide data on the long-term course of the disease, prognostic factors, and optimal immunotherapeutic regimes.

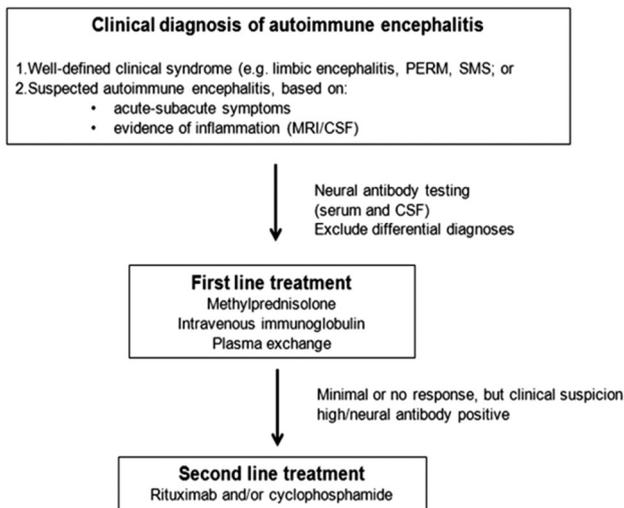


Figure 1: Suggested algorithm for approach to suspected autoimmune encephalitis.^[61-63] PERM: progressive encephalomyelitis with rigidity and myoclonus; SMS: stiff-man syndrome; MRI: magnetic resonance imaging; CSF: cerebrospinal fluid

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