A CASE OF INFLAMMATORY PERIPHERAL NERVE DESTRUCTION ANTEDATEING ANTI-NMDA RECEPTOR ENCEPHALITIS

NMAD receptor (NMDAR) antibody encephalitis is the second most common autoimmune encephalitis after acute disseminated encephalomyelitis, presenting with a multistage clinical progression with prodromal signs, psychiatric abnormalities, reduced levels of consciousness, epileptic seizures, dyskinesias, and autonomic dysfunction. Pathogenic NMDAR autoantibodies downregulate NMDARs in cultured hippocampal neurons and after injection into the rat brain. Teratomas are common in adult women and are believed to trigger the antibody response against NMDARs by ectopically expressing neuronal antigens. Causal factors in tumor-negative patients are largely unknown. Recent studies showed that infectious encephalitides with herpes simplex virus (HSE) are associated with the generation of NMDAR antibodies and that antibodies cause immunotherapy-responsive relapsing symptoms post-HSE. It is unknown, however, whether further types of neuronal damage can trigger a secondary immune response leading to NMDAR encephalitis.

Case report. A previously healthy 75-year-old man was referred with confusion, disorientation, and agitation. History revealed signs of neuropathy with ascending pain and numbness in his legs starting at least 3–4 weeks (but not >8 weeks) prior to referral; a friend reported progressive difficulties in walking and frequent falls during this time. Lumbar puncture showed normal cell count (5/μL) but high protein (>1,047 mg/L). Viral encephalitis was suspected and acyclovir was started until PCR for herpes simplex virus (HSV) and varicella-zoster virus showed negative results. Head MRI was unremarkable, showing mild generalized atrophy. No infectious agents were detected in serum and CSF, and NMDAR antibodies were below detection threshold (figure).

Encephalopathy progressed and the patient fluctuated between severe agitation with hyperkinetic movements and unresponsiveness, later developing unexplained fever and respiratory distress requiring mechanical ventilation. EEG showed bilateral slowing. Repeated serum and CSF testing showed increasing titers of immunoglobulin (Ig) G–NMDAR antibodies (figure). NMDAR antibody testing was performed using a standardized laboratory assay (figure); further antibodies against neural antigens (AMPAR, GABAβR, GlyR, LGI1, CASPR2, AQP4, DNER(Tr), GAD65, Hu, Yo, Ri, Ma/Ta, amphiphysin, MAG, myelin) were absent (Euroimmun, Lübeck, Germany).

CT with contrast enhancement and PET scan showed no evidence of a tumor; brain PET revealed bilateral hypometabolism in thalamus, caudate nucleus, and cerebellum. Immunosuppressive therapy was started with methylprednisolone and IVIg. Four cycles of plasma exchange and 2 doses of 1,000 mg rituximab were started after 6 and 12 weeks of limited improvement, respectively. Therapy resulted in marked improvement of leg pain, weakness, and NMDAR encephalitis, paralleled by loss of antibodies (figure).

Differential diagnosis of the concomitant severe neuropathy included nerve conduction studies, which showed sensorimotor axonal and demyelinating neuropathy predominantly affecting the legs (compound muscle action potential of right tibial nerve was reduced, nerve conduction velocity was delayed, and there were no reproducible recordings from peroneal and sural nerves). EMG was refused by the patient. Reflexes were normal, and there were no atrophies or fasciculations. Sural nerve biopsy 3 months after referral showed severe reduction of myelinated fibers but also strong axonal degeneration and single axon regeneration clusters (figure). Diagnosis of acute neuritis was established. There were no findings supporting vasculitis, paraproteinemia, vitamin B12 deficiency, diabetes, hepatitis C virus, HIV, or porphyria as alternative diagnoses.

Nine months after initial presentation he showed no dyskinesias, no signs of autonomic or psychiatric dysfunction, and normal consciousness and attention. Motor function and leg pain were markedly improved whereas electrophysiology was unchanged. Persisting anterograde memory deficits suggest postencephalitic residuum.

Discussion. The case describes how subacute onset of severe neuritis antedates classical NMDAR encephalitis. The temporal relationship of symptoms and antibody kinetics makes it unlikely that both diseases result from a shared precipitating factor, although such associations were shown for NMDAR encephalitis and demyelinating disorders. We hypothesize that NMDAR antibodies represent a secondary immune response to massive
destruction of peripheral nerve proteins as shown in sural nerve biopsy, also supported by the temporal relationship between neuritis onset and NMDAR antibodies. It is well known that peripheral nerves contain high amounts of NMDAR protein, which becomes phagocytosed after disintegration of the nerve. Transport to lymphoid organs may then trigger an immune response resulting in encephalitis.

In this way, the case adds a novel mechanism to the most relevant question: which pathophysiologic steps lead to NMDAR encephalitis? While teratomas have been shown to express NMDAR protein and are associated with NMDAR encephalitis, infectious encephalitides with HSV were recently added to the still-small list of etiologies. The present case suggests that inflammatory neuropathies may also trigger a secondary autoimmune response leading to severe encephalitis. Whether less obvious damage to peripheral neuronal tissue can under certain conditions trigger an immune response with formation of pathogenic antibodies is unknown.
autoantibodies and NMDAR encephalitis is an intriguing question.

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A case of inflammatory peripheral nerve destruction antedating anti-NMDA receptor encephalitis

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