Anti-NMDA Receptor Encephalitis in an Adolescent

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Background

Anti-NMDA receptor encephalitis is an autoimmune disease associated with antibodies against N-methyl-D-aspartate (NDMA) receptors, which are involved with synaptic transmission and plasticity. These autoantibodies can variably be associated with tumors and are responsive to immunotherapy.

We present a case of an adolescent male with a history of a pineal gland germinoma, found to have Anti-NMDA receptor encephalitis requiring methylprednisolone, intravenous immunoglobulin (IVIG), therapeutic plasma exchange, Rituximab, and Sirolimus therapies.

Case Report

We report a 17 year old male who presented with altered mental status and worsening hallucinations. Past medical history includes PTSD, ADHD, and pineal gland germinoma status post radiofrequency ablation.

Presentation was remarkable for slow, dysarthric speech, mental disorientation, and delayed movements upon simple commands. Otherwise, vitals, baseline labs, imaging (Brain MRI), and diagnostics (EEG) were unremarkable.

A multidisciplinary team decided to initially undergo psychiatric therapeutic intervention, focused on benzodiazepines and antipsychotics, however, patient’s status worsened, demonstrating catatonia, dystonia, and explosive agitation. Further laboratory investigation ruled out electrolyte disturbances, uremia/hepatic encephalopathy, thyroid storm, drug overdose/withdrawal, eventually settling on a working diagnosis of catatonic depression.

Given his deteriorating mental status throughout admission and prior oncologic history, a lumbar puncture was performed to rule out other causes of encephalopathy, with CSF studies remarkable for the presence of NMDA encephalitis autoantibodies, most likely sequelae from his existing pineal gland germinoma.

Initial treatment consisted of methylprednisolone, intravenous immunoglobulin (IVIG), and therapeutic plasma exchange. After minimal success, patient received rituximab and sirolimus, which resulted in improved psychomotor symptoms.

Psychiatric Diagnoses Reported in Anti-NMDA Receptor Encephalitis

- Psychosis including catatonia
- Schizophrenia
- Bipolar Disorder
- Depression
- Conversion Disorder
- ADHD
- PTSD

Discussion

Diagnosing Anti-NMDA receptor encephalitis can be quite challenging, particularly with its variable presentation in the setting of prior psychiatric diagnoses. Classically, Anti-NMDA receptor encephalitis demonstrates hyperkinetic movements (e.g. dystonia, dyskinesia, chorea, seizures); however, catatonic movements do occur, and can be a severe and life-threatening state. Being cognizant of the overlapping psychiatric and psychomotor symptoms, including catatonic features, can prompt the clinician to suspect and rule out Anti-NMDA encephalitis prior to settling on a primary psychiatric diagnosis.

References


Disclosure: Authors of this presentation have the following to disclose concerning possible financial or personal relationships with commercial entities that may have a direct or indirect interest in the subject matter of this presentation:

- Cynthia Carter – nothing to disclose
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