

complicated by a UTI and delirium. PLEX was restarted after one week and her mental status improved, reportedly 80% back to baseline after five sessions.

Conclusions

The main diagnostic challenge for LE is recognizing its unique hallmark, FBDS, often associated with insidious decline in cognition. FBDS may have no ictal correlate on EEG. Early diagnosis and management prevents long term disability - mainly cognitive. LE is commonly treated with immunotherapy, corticosteroids and PLEX. Monitoring response is challenging as patients are commonly older with multiple comorbidities: steroid-induced psychosis and delirium may complicate evaluation and treatment side effects limit options.

Disclosure: Dr. Sadeghpour has nothing to disclose. Dr. Neupane has nothing to disclose. Dr. Mouti has nothing to disclose. Dr. Clark has nothing to disclose.

Autoimmune Encephalitis-A Diagnostic Challenge for the Neurologist

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Objective

Autoimmune encephalitis, is a clinically challenging entity with varied neurological presentations. As autoimmune serology is negative in over 50% cases, our objective was to prioritise the use of electroencephalography supported by MRI Brain or PET-CT imaging to make a definitive diagnosis of autoimmune encephalitis.

Background

Autoimmune encephalitis is the consequence of an antibody mediated neuronal damage caused by cell surface antigens. However, antibody assays can be negative in early stages of the disease. Hence, we suggest the use of electroencephalogram along with MRI brain imaging or PET-CT scans to avoid diagnostic delays.

Design/Methods

During a span of four years [2018–2022] a retrospective review of the case records of 50 patients of autoimmune encephalitis and 50 patients of non-autoimmune encephalitis were compared. Besides clinical examination, serum and cerebrospinal fluid viral and autoimmune antibody assay, electroencephalogram, Magnetic Resonance and FDG-PET-CT scans were used to confirm the diagnosis.

Results

60% patients were seronegative and 40% were seropositive in the autoimmune group while 90% were seropositive and 10% seronegative in the non-autoimmune encephalitis group. Electroencephalography was abnormal in all cases of autoimmune encephalitis 100% [50/50] and in 80% [40/50] cases of viral encephalitis. In autoimmune encephalitis, MRI Brain revealed evidence of limbic encephalitis in 80% cases and FDG PET-CT scans were abnormal in the remaining 20%. In non-autoimmune encephalitis, MRI Brain was abnormal in 60% cases and FDG PET-CT was abnormal in 10%.

Conclusions

In seronegative autoimmune encephalitis, electroencephalographic abnormalities supported by MRI Brain imaging and PET-CT scans enabled early diagnosis in 100% cases [p value <0.001]. While in non-autoimmune

encephalitis serology with electroencephalography, MRI Brain and PET-CT scans were diagnostic in 80% cases [p value <0.001]. However, as the sample size is small further studies are needed to confirm these findings.

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Brains on Fire: Patient Outcomes and Quality of Life Following Autoimmune Encephalitis

Ava Easton

Objective

To present and improve understanding of patient outcomes and quality of life post-autoimmune encephalitis.

Background

Patient outcome following encephalitis and in particular following autoimmune encephalitis is not well understood. It is only over the last 15 years that we knew, became able to test for, and identify some of these autoimmune causes of encephalitis and so there currently only a small and emerging literature about patient outcomes. Many papers that talk about or refer to patient outcome look quite early on in the patient pathway and often when they talk about outcome they mean immediate clinical outcome and not how patients are, if they survive, several months or years down the line. Therefore we don't actually have a good handle yet on autoimmune patient outcomes nor their quality of life. Yet, most papers agree that much more needs to be done to assess long-term outcome and quality of life in autoimmune encephalitis patients. One further point of importance is that some complications of AE might not appear until several months or years down the line.

Design/Methods

Review of the literature accompanied by first-hand patient video testimony providing rich insight into long term outcomes and impact on quality of life post-autoimmune encephalitis.

Results

Patient outcomes post-AE can be life-changing and in some instances may occur several months and years post-acute illness. These outcomes and quality of life influence how patients understand and make sense of their experience; how they engage with recovery and rehabilitation. A range of poorly understood factors influence patient quality of life post-AE. These include a lack of easily understood index event, lack of community and collective understanding, fears of relapse, post-ICU PTSD, and a lack of accurate information and understanding of their outcomes pre and post-discharge.

Conclusions

Improved understanding of patient outcomes post autoimmune encephalitis can improve patient care and engagement.

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