Complete right cerebral hemispheric diffusion restriction and its follow-up in a case of Rasmussen’s encephalitis

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Accepted 23 August 2015

DESCRIPTION
A 30-year-old man presented with generalised status epilepticus. Initial brain CT was normal (figure 1). His generalised seizures were controlled with antiepileptics. However, he continued to have left focal seizures. An MRI was performed showing diffuse T2 hyperintensity, oedema and high signal on diffusion-weighted imaging (DWI) with corresponding low signal on apparent diffusion coefficient (ADC) maps in the entire right cerebral hemisphere, implying restricted diffusion (figure 2A–C). Cerebrospinal fluid (CSF) analysis showed raised protein at 70 mg/dL and increased lymphocytes in cell count. CSF PCR for herpes simplex virus was negative. A working diagnosis of hemisphere demyelination was performed. The patient responded to intravenous methylprednisolone and was continued on oral steroids for 2 months. He was able to walk at the time of discharge with a power of 4 plus on the left.

Five months later he returned in a state of unconsciousness, with left focal seizures. He also had a fever and productive cough. He had discontinued the antiepileptic drugs. A diagnosis of left focal status epilepticus with aspiration pneumonia and encephalopathy was performed. EEG showed slowing with sharp waves in the right cerebral hemisphere (figure 3). MRI showed diffuse right cerebral hemiatrophy with gliosis and high signal on ADC maps, suggestive of facilitated diffusion (figure 4A–C). The imaging findings along with the clinical presentation and progression clinched the diagnosis of Rasmussen’s encephalitis. The patient was treated with broad spectrum antibiotics and antiepileptic drugs. His sensorium improved but he had residual left hemiparesis on discharge with a power of 3/5.

Rasmussen’s encephalitis is a chronic progressive inflammatory disease of unknown aetiology, usually involving one cerebral hemisphere. Clinically, patients have abrupt onset of seizures with progressive neurological decline.1–3 Diagnosis can be performed on the basis of the clinical picture and neuroimaging demonstrating progressive cerebral hemiatrophy.1 Imaging in the acute phase may

![Figure 1](image1.png) Normal brain CT at initial presentation.

![Figure 2](image2.png) MRI in the acute phase of illness, taken 4 days after the brain CT study. (A) Axial T2-weighted image showing diffuse hyperintensity and oedema of the right cerebral hemisphere. (B) Diffusion-weighted image showing diffuse hyperintensity in the right cerebral hemisphere. (C) Apparent diffusion coefficient (ADC) image showing diffuse hypointensity noted in the right cerebral hemisphere; findings in B and C indicate diffusion restriction.
reveal cerebral oedema. Cerebral atrophy, gliosis and facilitated diffusion are the classical features described with progression of the disease. These features were demonstrated in our case. However, complete right cerebral hemispheric diffusion restriction seen in the acute phase in our patient has not been previously reported, to the best of our knowledge. We aim to highlight this finding and offer an explanation for it.

DWI is a MRI technique that reflects the ability of water molecules to move freely in tissue. It reveals the molecular state of the tissue rather than giving a specific diagnosis. Restricted diffusion in an area implies that the motion of water molecules in that area is impeded as compared to normal. Restricted diffusion is seen as hyperintensity on DWI images, with corresponding area of hypointensity on ADC images. On the other hand, facilitated diffusion/increased diffusivity implies there is increased movement of water molecules. On imaging, it is seen as a variable signal intensity on DWI and increased signal intensity on ADC images. In our case, the finding of restricted diffusion in the acute stage can be explained on the basis of cytotoxic oedema, which impedes the free movement of water molecules. The finding of facilitated diffusion in the follow-up images is explained on the basis of atrophy and gliosis in the chronic stage of Rasmussen’s encephalitis, which corresponds with the study of Sener.

**Learning points**

- Rasmussen’s encephalitis is a chronic progressive inflammatory disease of unknown aetiology, usually involving one cerebral hemisphere. Clinically, patients present with seizures and progressive neurological decline.
- Diagnosis can be performed on the basis of the clinical picture and neuroimaging. The finding of unilateral hemiatrophy in the appropriate clinical setting clinches the diagnosis.
- Diffusion restriction on MRI is typically associated with diagnosis of an acute infarct, however, it can also be seen in various other conditions. In our case, we attribute diffusion restriction to cytotoxic oedema in the acute phase of the illness, before the onset of atrophy.
- In the later phase, diffusion is facilitated, which corresponds with atrophy and gliosis.
Contributors FF contributed to the concept, collection of data and drafting of the article. AB helped in collection of data and in drafting the article. ST was the clinician in charge of the patient. He contributed to the diagnosis and management of the patient and reviewed the paper prior to submission. RV overlooked the drafting of the paper and reviewed the paper prior to submission. All the authors contributed to the diagnosis and reviewed the paper prior to submission.

Competing interests None declared.

Patient consent Obtained.

Provenance and peer review Not commissioned; externally peer reviewed.

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