Eclampsia with Posterior Reversible Encephalopathy Syndrome

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Abstract

Background: The posterior reversible encephalopathy syndrome (PRES) or reversible posterior leukoencephalopathy syndrome (RPLS) includes headache, visual disturbance, and seizure, altered mental status with or without focal neurological signs. Here, we present a 24 year old postpartum female with PRES in the setting of post-partum eclampsia with cortical blindness and seizure. Case Report: A 24 year old African American female presented with severe, constant, throbbing, bilateral occipital headache nine days after normal spontaneous vaginal delivery followed by blurred vision for two days. It was complicated by generalized tonic-clonic witnessed seizure. Magnetic Resonance Imaging (brain) showed findings consistent with posterior reversible encephalopathy syndrome. The patient was successfully managed with intravenous labetolol and magnesium sulfate for eclampsia in intensive care unit. Conclusion: Focussed clinical approach, early detection of significant neuroimaging findings upon presentation, timely aggressive treatment and cessation of causal agents are crucial for satisfactory recovery and outcomes. This syndrome is overlooked by the physicians and usually radiologists are the saviors for timely diagnosis for PRES.

Keywords: Eclampsia, Magnetic Resonance Imaging, Posterior Leukoencephalopathy Syndrome, Pregnancy, Seizures.
sinus bradycardia at 50 bpm. Trans-esophageal echocardiogram showed normal ejection fraction and no atrial septum defect or ventricular septum defect. Computed Tomography scan (head) without contrast [Fig.1] showed normal and repeated CT scan [Fig.2] showed no evidence of acute intracranial hemorrhage, fluid collections or mass effect. Magnetic Resonance Imaging (brain) [Fig.3,4] showed cortical and sub-cortical regions with abnormal FLAIR (Fluid Attenuated Inversion Recovery) signal with associated bright diffusion signal involving the frontal and parietal lobes as well as the occipital lobes and left cerebellar hemisphere consistent with posterior reversible encephalopathy syndrome (PRES). The eclampsia was managed with intravenous labetolol and magnesium sulfate with prophylactic levetiracetam. Next morning, her vision improved but still had difficulty seeing the features of a person’s face. She was discharged home with instructions to follow up with neurologist.

Discussion

The posterior reversible encephalopathy syndrome (PRES) or reversible posterior leukoencephalopathy syndrome (RPLS) is usually neuro-imaging finding of reversible vasogenic subcortical tissue swelling without gross infarction. The underlying patho-physiological hypothesis is (i) cerebral ischemia from vasospasm (ii) disordered cerebral autoregulation and (iii) endothelial dysfunction [1,2]. The incidence is common in young than elderly and in female than male. It is usually presents with clinical seizure in 87%, headache unresponsive to medications in 53%, encephalopathy in 92% and visual disturbances in 39% [3].

Typically, Computed Tomography (CT) scan is less significant in PRES but Magnetic Resonance Imaging (MRI) of the brain demonstrates symmetric, posterior occipito-parieto-temporal area white matter edema. However, they may present with atypical locations or asymmetric lesions. The involvement of frontal lesions is common although

Fig.1: CT scan without contrast (on admission) showed normal the ventricles, sulci and cisterns without evidence of hydrocephalus.

Fig.2: CT scan without contrast (repeated) showed no suspicious interval change.

Fig.3: MRI without contrast showed cortical and sub-cortical regions with abnormal FLAIR signal with associated bright diffusion signal (white arrows) involving the frontal, parietal lobes, the occipital lobes and left cerebellar hemisphere consistent with posterior reversible encephalopathy syndrome.
the lesions may be detected in the brainstem and basal ganglia [4]. MRI showed increased T2 and FLAIR (Fluid Attenuated Inversion Recovery) signal due to cerebral edema in the posterior region of the cerebral hemisphere especially the posterior parieto-occipital lobes as those areas are less sympathetic innervated. The increased signal is usually bilateral and symmetrical but sometimes it may be single and unilateral [5]. In approximately 56% of the patients, the regions of vertebra-basilar territory such as thalamus and cerebellum are involved [6]. Increased T2 and DWI (diffusion weighted imaging) signal seen as water content in vasogenic edema is high. In setting of vasogenic edema in PRL, DWI is decreased while mobility is increased but in cytotoxic cerebral edema, DWI is increased but water mobility is decreased owing to cerebral edema.

Whenever PRES is in doubt, management should be aggressive as delayed treatment might end up with permanent brain damage. Treatment includes proper blood pressure control, cessation of causal agent or therapeutic agent; manage underlying co-morbidities and seizure prophylaxis. In patient with hypertensive encephalopathy, mean arterial pressure must reduce 20-25% in first two hours and diastolic pressure to 100 mmHg as rapid reduction in blood pressure worsens end organ perfusion and cerebral function.

**Conclusion**

In our case of PRES due to eclampsia, magnesium sulfate is preferred agent for prophylactic seizure owing to neuro-protective effects by stabilizing of cerebral circulation and stabilization of neuronal membranes and inhibiting in stimulatory neuron transmitters, such as glutamate [7,8] and magnesium is superior to diazepam and phenytoin. For PRES due to pre-eclampsia, the mainstay is delivery of placenta and fetus. Most PRES patients will have complete neurological recovery usually in two weeks as in our patient and seizure is in remission once neuroimaging changes have recovered in three to seven days [9]. Complete reversibility is pathognomonic features of PRES or RPLS. However, some cases can progress to leukomalacia and permanent cerebral ischemia and damage particularly in setting of PRES due to immunotherapeutic agents [10].

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