Case Report

PRES in pregnancy: MRI and it's role in decision making

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ABSTRACT

Posterior reversible encephalopathy syndrome (PRES) is an alarming clinic-neuro-radiological syndrome accompanying various clinical conditions, presenting with headache, encephalopathy, seizures, cortical visual disturbances or blindness. The lesions in PRES are thought to be due to vasogenic oedema, predominantly in the water-shed regions of the posterior cerebral hemisphere, fortunately, completely reversible with management of the primary condition. We report a case of primigravida with 26 weeks pregnancy, who presented with acute, severe, intractable, throbbing headache, tingling sensation in the posterior neck and upper shoulder region, photophobia, nausea and mild pre-eclampsia. Standard therapy did not relieve the headache, BP remained fluctuant, so migraine/some intra-cranial pathology was suspected. MRI is the gold-standard diagnostic modality. It revealed the classical acute PRES picture. PRES implies breaching of the blood brain barrier, resultant cerebral edema and potential for further intra-cranial events of serious proportions. Notably, PRES may be seen with normotension. Pregnancy was terminated un-eventfully, recovery was prompt and complete. Two weeks later, the patients showed marked improvement clinically and neuro-imaging features of PRES had dis-appeared.

Keywords: PRES, Pre-eclampsia, Cerebral edema, MRI, Pregnancy

INTRODUCTION

Posterior Reversible Encephalopathy Syndrome (PRES) refers to a rare clinico-radiologic entity introduced as late as 1996 by Hinchey et al., where a neurotoxic state is coupled with a unique CT or MRI picture. Recognised in the setting of a number of complex conditions including pre-eclampsia/eclampsia, it has characteristic features on neuro-imaging and non-specific symptoms comprising headache, confusion, encephalopathy, cortical visual disturbances or blindness and seizures.2,3

CASE REPORT

We report a case of primigravida with 26 weeks pregnancy, presenting with acute, severe, in-tractable,
Symptoms dis-appeared with remarkable speed and neuro-imaging features of PRES reversed miraculously.

**Figure 1:** MRI showing classical picture of acute PRES- T2 weighted images showing hyperintensities in occipital and parietal lobes characteristic of posterior reversible encephalopathy syndrome.

**DISCUSSION**

Posterior reversible encephalopathy syndrome, also termed reversible posterior leukoencephalopathy syndrome or posterior cerebral oedema syndrome is a newly recognized clinico-neuro-radiological syndrome, affecting predominantly the white matter of the posterior cerebral hemispheres.

PRES is a grave complication accompanying a myriad of obstetric and medical conditions, chiefly, hypertension, preeclampsia/eclampsia and HELLP syndrome. Others being hypertensive encephalopathy, nephropathy, following immunosuppressive or anticancer medication (methylprednisolone, cyclosporine and tacrolimus), systemic lupus, amyloid angiopathy, Henoch-Schonlein syndrome, hemolytic uremic syndrome, sepsis (gram positive), shock. Headache is typically acute, severe, intractable, associated with visual disturbances, seizures, altered consciousness, nausea, vomiting, focal neurologic deficits, stupor and coma, all indicative of multi-focal intra-cranial events.⁴⁻⁵

Abrupt increase in the blood pressure, leading to an acute disruption of the blood-brain barrier, causing the brain to swell is the triggering event. Parietal and occipital lobes are most commonly involved, followed in-frequently by frontal lobes, the inferior temporal- occipital junction, and the cerebellum.⁶ Focal patchy areas of vasogenic edema due to PRES may also be seen in the basal ganglia, brainstem, deep white matter and internal capsule.⁷ Myogenic and neurogenic component form the cerebral auto-regulatory mechanism which maintains constant brain perfusion. Effectiveness of neurogenic component is directly proportional to degree of sympathetic innervation. To maintain constant perfusion, vasocostriction occurs in response to hypertension via sympathetic innervations of the cerebral vessels. Sudden increase in BP, overcomes auto-regulatory capability of brain vasculature leading to abrupt dilatation of cerebral arterioles. Hyper-perfusion ensues. Breakdown of blood brain barrier causes petechial haemorrhages. Focal transudation of fluid into interstitium results in vasogenic edema. Posterior circulation is thought to be more susceptible to this type of damage, because there is less sympathetic innervation of the vertebrobasilar vasculature to protect the parenchyma from rapid increases in arterial blood pressure.¹¹⁻¹²

On MRI, T1-weighted images show hypo-intense and T2-weighted images show hyper-intense areas.⁸ The typical imaging findings of PRES are most apparent as hyper-intensity on FLAIR images in the parieto-occipital and posterior frontal cortical and subcortical white matter, less commonly brain stem, basal ganglia and cerebellum are involved. Atypical imaging appearances include contrast enhancement, hemorrhage and restricted diffusion on MRI.⁹ PRES lesions involving the occipital lobe spare the calcarine and para-median occipital lobe. This feature, along with the predominant involvement of white matter helps to distinguish this syndrome from bilateral infarctions of the posterior cerebral artery.¹ When regions of the brain other than the parieto-occipital lobes are predominantly involved, the syndrome can be called atypical. In such cases, a diffusion weighted MRI with ADC (increased apparent diffusion coefficient) mapping shows increased ADC values representing vasogenic edema in these areas, thus differentiating atypical PRES from other brain disorders.¹⁰ Diffusion-weighted imaging may also help distinguish vasogenic edema (increased apparent diffusion coefficient) from cytotoxic edema (reduced apparent diffusion coefficient), seen in acute arterial ischemic injury. Magnetic resonance angiography using 3D Time of flight (TOF) technique, reversible “vasculopathy” (diffuse/focal vasocostriction) or vessel pruning is noted.⁵

Treatment of elevated blood pressure is considered central in the management of PRES. Generally, the goal of treatment is the reduction of mean blood pressure to premorbid levels. Aggressive intravenous antihypertensive therapy should commence immediately, BP to be closely monitored, with an arterial catheter. IV Nicardipine (5-15 mg/h) and Labetalol (2-3 mg/min) are considered first-line medications in PRES.¹³

When the under-lying cause of PRES is treated effectively, MRI lesions shows resolution, unless the condition unfortunately, progresses to infarction or hemorrhage.¹

**CONCLUSION**

Strong clinical suspicion and MRI, being the gold-standard, helped us to arrive at the diagnosis of PRES, a serious and potentially lethal condition. Further, we could distinguish PRES from other neurological conditions. MRI helped in the prompt diagnosis and early intervention. The disease was halted in the reversible stage, MRI, then, is the best available diagnostic modality in patients with cerebral signs and symptoms complicating pregnancy further permitting early intervention.
REFERENCES


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