Posterior Reversible Encephalopathy Syndrome (PRES)
(Report of three cases with brief review of literature)

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Abstract: We report three cases of Posterior Reversible Encephalopathy Syndrome in different settings. First case was associated with systemic lupus erythematosus with gestational hypertension and lupus nephritis. Second case had no history of hypertension while third patient had only transient rise of blood pressure during & after an episode of seizures. All three cases had complete recovery. Posterior Reversible Encephalopathy Syndrome is a clinical-neuroradiological term, often associated with delayed postpartum eclampsia & classical imaging features usually in the form of bilateral, symmetrical, reversible white matter oedema involving commonly the posterior cerebral circulation. Diagnosing these lesions early & providing appropriate treatment is important to achieve an optimal neurological outcome.

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I. Introduction
Most cases of eclampsia usually occurs from 20 weeks of gestation up to first 2 days after delivery¹. Some cases of eclampsia have been reported as late as 23 days postpartum². In classical case of eclampsia, the pre-eclamptic prodrome of proteinuria and hypertension precedes the onset of generalised seizure¹.
In contrast, Delayed postpartum eclampsia is characterised by the onset of generalized seizures in a puerperal women from 2 days up to 30 days postpartum, frequently in a patient who had a normal pregnancy and delivery and without pre-eclamptic prodrome of proteinuria & hypertension³,⁴. Posterior Reversible Encephalopathy Syndrome is an obstetric emergency frequently occurring in a postpartum period.

II. Case History

Case 1:
A 25 years old patient gravida 3, para 1, living 1, abortion 1 had history of 9 months amenorrhea was admitted with chief complaints of swelling over the both feet since 2 months and oral ulcers since last 20 to 25 days. Patient was a diagnosed case of Systemic Lupus Erythematosus (SLE) with Gestational Hypertension since 6 months. On physical examination, the patient was afebrile with tachycardia of 104/min, blood pressure was 160/100 mmHg, bilateral pitting pedal oedema was present. Urine dipstick showed 4+ albuminuria.

In view of high risk pregnancy (previous Lower Segment Caesarean Section, Gestational hypertension & Lupus Nephritis), patient was taken for elective LSCS. LSCS was uneventful with delivery of healthy baby.

On 12th day postpartum patient experienced intense headache, nausea, vomiting and blurring of vision, followed by generalized tonic-clonic seizures. The aetiology of seizures was unclear initially. The differential diagnosis included subarachnoid haemorrhage, space occupying lesion in brain, intracranial haemorrhage, ischemic stroke, venous thrombosis, HELLP syndrome, Meningo-encephalitis.

Computed Tomography (CT) Scan of brain revealed subcortical white matter hypodensities in bilateral occipital lobes, mostly representing vasogenic oedema suggest possibilities of posterior reversible encephalopathy. MRI and MRV were ordered to rule out thrombosis and space occupying lesion. MRI scan of brain showed gyral oedema in bilateral frontal and parieto-occipital parenchyma suggest possibilities of reversible encephalopathy syndrome. MRV was normal. The diagnosis of PRES was confirmed by imaging and clinical course as well as by ruling out other aetiologies. Patient given antihypertensives and magnesium sulphate and anti-epileptic. The patient had no further seizures, recovered slowly and was discharged on hospital day 40.

Case 2:
A 23 years old patient gravida 1, para 1 had normal pregnancy and normal full term vaginal delivery of healthy baby and discharged from hospital on 3rd day postpartum. On 4th day postpartum patient experienced mild headache, nausea and blurred vision in morning hours to which patient neglected. On the same day

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afternoon patient developed generalised tonic-clonic seizures for which she was admitted. On physical
examination patient was afebrile, blood pressure was 130/80 mmHg. Patient’s serial blood pressure recordings
were within normal limit. Urine dipstick showed no albuminuria. Patients all the laboratory investigations and
metabolic profile were within normal limit. Patients MRI brain with MRV & MRA were ordered which showed
areas of altered signal seen in both occipito-parietal and frontal regions dominantly involving the junctional
zone of cortex and white matter with adjacent focal oedema and early ischemia seen suggest possibility of
posterior reversible encephalopathy. MR Angiography & venography revealed normal report. Patient given
magnesium sulfate and improved.

Case 3:
A 24 years old patient primigravida had normal pregnancy, underwent LSCS in view of prolonged
labour and foetal distress. LSCS was uneventful with delivery of healthy baby. On 6th postoperative day patient
developed intense headache, vomiting followed by generalised tonic-clonic seizures. At the time of seizures &
after short period of seizures her blood pressure was 170/100. Her ANC follow up record showed that patient
was previously healthy and normotensive. Her haemogram, serum uric acid, urine examination & fundus
examination was normal. MRI Brain showed bilateral symmetric fronto-parietal cortical abnormalities with mild
swelling, results consistent with PRES. MRA & MRV were normal. Patient given antihypertensive for short
period and patient was discharged on hospital day 12.

III. Discussion

PRES was first described by Hinchey & colleagues in 1996 as Reversible Posterior Leukoencephalopathy Syndrome. PRES has been described as various non-specific symptoms of intense headache, visual problems (e.g. blurring of vision, hemianopia, cortical blindness, hemineglect), altered consciousness and generalised seizures. Rarely the patients may develop focal neurological deficits like paresis.

PRES may develop suddenly or over several days. The term PRES is a misnomer as it occurs not always in posterior cerebral circulation but may occur primarily in anterior cerebral circulation. Also, PRES is not always reversible; it can be irreversible if the aetiology is not treated. In addition, patient in PRES may not always present with an encephalopathy, instead patient may have various nonspecific clinical features other than encephalopathy. In PRES white matter is mostly involved but it may involve deep white and gray matters.

Risk factors:
PRES a recently described clinical-neuroradiological term that is associated with several medical
condition besides preeclampsia/eclampsia and hypertension e.g. Renal failure, Post-transplantation (Allogeneic
bone marrow transplantation; Solid organ transplantation), Immunosuppressive therapy (Cyclosporine;
Tacrolimus) Autoimmune diseases (SLE; Systemic Sclerosis; Wegener’s Granulomatosis; Poly Arteritis
Nodos), Post-cancer chemotherapy and has recently shown to be associated with infection, sepsis, and shock.

Radiological findings of PRES:
The characteristic imaging in PRES is cortical or subcortical areas of hypoattenuation with a
predominantly bilateral, symmetrical, posterior distribution in the parietal & occipital white matter on CT or T2
hyperintensity on MR imaging. On MR imaging, the findings are most apparent on fluid-attenuated inversion
recovery (FLAIR) images. Most evident on T2-weighted MRI images, the lesions are hyperintense and located
at the gray-white junction. The parietal and occipital lobes are most commonly affected, followed by in
descending order the frontal lobes, the inferior temporal-occipital junction, and the cerebellum. Involvement of
the basal ganglia, brain stem, and deep white matter including the splenium of corpus callosum can also be
seen. Frequent involvement of posterior cerebral circulation is not well understood. It is presumed that
intracranial arterioles in anterior circulation have extensive sympathetic innervations than in posterior
circulation which are responsible for protection from raised blood pressure. This shows that the amount of
sympathetic innervation and the degree of parenchymal involvement are inversely proportional in PRES.

However, in some cases the protective effect of sympathetic innervation in anterior circulation can be
overcome and anterior circulation can be involved called as Atypical PRES. McKinny & colleagues reported
that PRES can be unilateral. This atypical unilateral involvement can cause difficulty in the diagnosis of PRES.
Follow up neuroimaging studies in PRES shows nearly complete resolution of imaging findings, and suggest
that PRES is usually associated with reversible cerebral oedema without infarction & confirms the diagnosis.

Pathophysiology of PRES:
The pathophysiology of PRES is not well understood. Vasogenic theory i.e. hypertension with loss of
autoregulation remains a widely accepted consideration for the development of brain oedema. Sometimes
patients with normal blood pressure may develop PRES if they have substantial rise in blood pressure which is
considered to be within the range of normal blood pressure. This is believed to be due to some neurotoxic
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PRES is seen in the absence of hypertension in 20%–40% of patients. Alternatively, endothelial dysfunction/injury, hypoperfusion, and vasoconstriction may lead to altered integrity of the blood-brain barrier. In the latter, though some degree of hypertension is present, reported blood pressure usually do not reach the limit of autoregulation (mean arterial pressure >160 mm Hg).

Treatment:
The objective of the treatment of PRES is directed at the underlying aetiology e.g., control of blood pressure, reducing the dose or withholding the offending drug in patients undergoing chemotherapy or immunosuppression. Some physicians use antiepileptic drugs or Magnesium sulphate to avoid the progress of seizures, but their role is controversial or yet to be established.

Prognosis:
The clinical outcome is variable but is mostly favourable with prompt treatment of the underlying cause and repeat neuroimaging may not be necessary. Immediate action to identify potential triggering drugs, controlling hypertension, and treating aetiology of PRES can lead to complete reversal of radiological and neurological findings. However, in few patients, PRES progresses to ischemia, infarction, or death.

IV. Conclusion
The pathophysiology responsible for the PRES remains controversial. PRES is usually reversible with early treatment of the aetiology responsible for it otherwise it has been shown to progress from reversible cerebral oedema to irreversible ischemic changes if appropriate treatment not started early. Early diagnosis and treatment is essential to ensure the best possible outcome. The occurrence of delayed postpartum eclampsia without typical preeclamptic prodrome of proteinuria & hypertension emphasizes the need of awareness of the diagnosis of PRES. Patients should be routinely counselled about the warning signs of PRES during postpartum period & on hospital discharge.

![Pathophysiology of PRES](image_url)

Fig. 1 Pathophysiology of PRES
REFERENCES