Blindness in eclampsia: CT and MR imaging

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SUMMARY Three cases of cortical blindness complicating eclampsia are described, with magnetic resonance imaging (MRI) and X-ray computed tomography (CT). The correspondence of MRI lesions (hyperintense on T2 weighted, and hypointense on T1 weighted sections) and low attenuation lesions on CT scan indicated ischaemia rather than haemorrhage as the pathological mechanism.

Reversible focal neurological lesions are a rare feature of eclampsia, most reports being of patients with cortical blindness. Where X-ray computed tomography (CT) has been carried out, this has either shown low attenuation lesions in the occipital cortex,1-7-9-11 multiple low attenuation lesions,6,7 or has been normal.12-15 We present three cases of blindness complicating eclampsia, and discuss CT and magnetic resonance imaging (MRI) findings.

Methods

Unenhanced CT was carried out using a Philips 310 Tomoscan, producing 6 mm axial cuts parallel to the orbito-meatal line. Corresponding T2 weighted (SE 2000/120) and T1 weighted (IR 1600/400/40) 8 mm thick axial MRI sections were obtained using a Picker 0.15 Tesla resistive imager.

Case 1

A 38 year old para 1 + 0 in the 38th week of pregnancy was admitted to hospital with a history of acute visual loss and frontal headache. She had been hypertensive during her first pregnancy and had no past history of migraine. The visual loss had woken her in the morning, and had progressed over two hours until she was blind. On admission to hospital, she was hypertensive with a blood pressure of 165/120 mm Hg, and proteinuric. She had a tonic/clonic seizure, eclampsia was diagnosed and a caesarean section was performed.

On transfer to the Regional Neurological Unit, she was drowsy, but oriented. She could perceive only strong light, but pupillary responses were normal. There was a conjugate palsy of upward and left lateral gaze. Tendon reflexes were exaggerated with clonus in the left leg. Plantars were downgoing. Vibration sense was absent below the waist, but sensory testing was otherwise normal. There were no rashes or other clinical evidence of systemic vasculitis. Blood pressure was 138/95 mm Hg.

ESR was raised to 86 mm in the first hour, she had a platelet count of 137 × 10^9/mm^3, and a urea of 7.4 mmol/l. Fibrin degradation products, and a clotting screen were normal. Autoantibody studies were negative. Cranial CT performed on admission showed no abnormality.

Her vision and abnormal neurological signs recovered to normal over 24 hours, despite continuing poor control of her blood pressure. MRI was performed on day 3 and showed...
focal lesions in both occipital poles. These were non-space occupying and gave a hypointense signal on the T1 weighted images, and a hypointense signal on the T2 weighted images (fig 1). Follow up MRI performed 2 months later showed complete resolution of the occipital lesions.

Case 2
A 20 year old para 0+0 had a caesarean section at 34 weeks for severe pre-eclampsia, diagnosed on the basis of hypertension, proteinuria and headache. She had no past history of hypertension or migraine. Post partum her blood pressure remained high despite treatment with atenolol, intermittently reaching levels as high as 200/100 mmHg. The fourth evening post partum she became confused and complained of severe frontal headache. The next morning she was alert and oriented, but her headache persisted. Over three hours her vision deteriorated until she was blind. She was transferred to the Regional Neurological Unit, where she suffered 2 tonic/clonic seizures. On examination, blood pressure was 165/110 mmHg. The fundi were normal, as were eye movements and pupillary reflexes. Vision had improved to perception of bright light throughout the visual fields. Neurological examination was otherwise normal.

Routine haematological and biochemical tests were normal. Cranial CT on admission showed bilateral occipital low attenuation lesions (fig 2). MRI carried out on the day of admission showed several focal non-space occupying lesions, hyperintense and hypointense on the T1 weighted and T2 weighted images respectively. These lay in both occipital lobes in the right posterior temporal cortex (figs 3 and 4), and in the white matter of the left temporal and occipital lobes, extending into the lateral cortex.

Vision returned to normal within 36 hours, despite her blood pressure remaining poorly controlled. Follow up MRI carried out 6 weeks later showed complete resolution of the occipital lesions, and near complete resolution of the temporal lesion.

Case 3
A 26 year old para 1+0 was admitted at 18 weeks gestation. Her previous pregnancy had been unremarkable, but ultrasound examination in this pregnancy carried out at 14 weeks had suggested foetal abnormality. She had been well until a few hours prior to admission when she had two tonic/clonic convulsions. She regained consciousness but then developed complete blindness over two hours. There was no history of migraine. On examination, visual acuity was limited to the perception of bright light. Fundal examination and pupillary responses were normal, as was the remainder of the neurological examination. Blood pressure was 150/90 mmHg, and she had proteinuria. Eclampsia was diagnosed, and the pregnancy terminated the same day.

Routine haematological and biochemical tests were normal, as was cranial CT performed on the day of admission. MRI performed on day 3, at which time visual acuity had recovered to counting fingers, showed a focal non-space occupying lesion in the right occipital pole, hyperintense and hypointense on the T1 weighted and T2 weighted images respectively (fig 5). Necropsy on the foetus revealed holoprosencephaly and a molar placenta.
Discussion

In all three cases the diagnosis of eclampsia seems certain, with hypertension, proteinuria and seizures occurring in association with pregnancy. Although Case 3 occurred at 18 weeks, this was in association with molar degeneration, an established risk factor for early eclampsia. Despite the statistical association between the two conditions, none of our patients had a history of migraine.

Blindness due to eclampsia may be due to lesions at any site along the visual pathway, but the majority of case reports are of cortical blindness. Although there were minor retinal changes in Case 1, intact pupillary reflexes and the radiological finding of occipital cortical lesions in our cases confirm that they too were of cortical origin. The majority of patients with cortical blindness due to eclampsia recover vision, over a period varying from 2 hours to 21 days, although there is one reported case of persistent deficit. In Case 1 blindness was the presenting feature of eclampsia, while in Case 2 it occurred 5 days post partum. Approximately 50% of the cases in the literature have occurred post partum, by up to 7 days. Blindness was the presenting feature of eclampsia in six cases.

Clinically, there also were lesions at other sites in Case 1. The finding of gaze palsies indicated a low midbrain/high pontine lesion. The sensory level to vibration sense suggested the possibility of a spinal cord lesion, as did the findings of increased reflexes and clonus. In the other two cases there was no clinical evidence of focal lesions other than those in the occipital cortex. In cases in the literature, increased tendon reflexes are common, and are not always associated with seizures. More definite evidence of pyramidal dysfunction such as ankle clonus, upgoing plantars and hemiparesis is also found as are gaze palsies, nystagmus and fluent dysphasias. In one case cortical lesions on CT coexisted with bilateral retinal detachments.

Most recent reports feature the results of CT, and this either was normal, showed focal low attenuation occipital lesions or more widespread low attenuation lesions. The lesions were non-enhancing, and were ascribed to localised oedema or to infarction. Repeat CT has shown partial resolution of lesions in 3–5 days and complete resolution in 9–14 days. A recent report describes the results of MRI in a single case of eclampsia. Although clinically the patient had no focal neurological features, lesions with similar signal characteristics to those seen in our cases were shown in the right occipital lobe and left parietal lobe. They had resolved by the time a further scan was carried out 3 weeks post partum, and were ascribed to localised oedema. They were not detected by CT. CT and MRI findings in hypertensive encephalopathy are similar.

Our MRI findings are consistent with those in
transient ischaemic lesions, and reflect focal increases in brain water content, which may persist for several days after the resolution of neurological signs. In contrast, acute focal haemorrhage gives a hypointense signal on T₁ weighted images and hyperintensity on T₂ weighted images. In the large necropsy series of Sheehan and Lynch and Govan gross haemorrhage was the most frequent cause of death, but the most common finding was of cortical groups of small (0.3–1 mm) haemorrhages associated with infarcts of similar size. It seems likely that our findings represent such lesions, and that they are primarily ischaemic rather than haemorrhagic. Small haemorrhages are seen in the placental vascular bed in eclampsia, and ultrastructural studies suggest that they are secondary to ischaemic vascular damage.

Angiographic studies in eclampsia show generalised large vessel spasm, and tend to support the view that ischaemia is the cause of focal neurological dysfunction. However, angiographically demonstrated spasm does not always relate well to cerebral blood flow, and spasm may occur without ischaemia and vice versa.

The finding of ischaemic lesions in eclampsia suggests that aggressive treatment of hypertension is likely to exacerbate neurological damage. It would appear that the most appropriate treatment in such cases would be a drug with cerebral vasodilator activity such as nifedipine. Alternatively, a more specific cerebral vasodilator agent such as nimodipine might be used in conjunction with other antihypertensive agents.

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