Reversible cerebral vasoconstriction syndrome in a postpartum female complicated by subarachnoid haemorrhage

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DESCRIPTION

A 31-year-old woman experienced sudden onset parieto-occipital headache 2 days postpartum. This resolved spontaneously but recurred at 6 days postpartum, when she reported the worst occipital headache she had ever experienced. Neurological examination was unremarkable. A non-contrast CT scan of the brain revealed subarachnoid haemorrhage (SAH) predominantly distributed...
The patient was urgently transferred to a neurosurgical unit and underwent cerebral digital subtraction angiogram which demonstrated multiple areas of arterial stenosis and dilatation, suggestive of diffuse vasospasm (figure 2). Blood tests for vasculitis and autoimmune disease, urine toxicology screen and cerebrospinal fluid analysis were normal.

Reversible cerebral vasoconstriction syndrome is characterised by sudden severe headache with cerebral arterial vasoconstriction which spontaneously resolves. It is more prevalent in females and around two thirds of cases are secondary to pregnancy or vasoactive substances including cannabis, cocaine, ecstasy, amphetamines and binge drinking. Patients are said to complain of an intense sudden onset headache which may relapse and remit as here. Neurological deficits and seizures may occur at presentation. A quarter to a third will have raised blood pressure during the episode.

Investigations are often directed to excluding aneurysmal subarachnoid bleed and non-contrast CT may show SAH in 20%, usually cortical. Infarction and intracerebral haemorrhage are less common at around 5%. Angiogram is definitive with the so-called ‘string of beads’ finding of segmental narrowing and dilatation as illustrated here. Nimodipine (60 mg six times daily), which was given here, is said to speed resolution, though more recently aggressive endovascular interventional techniques such as intra-arterial vasodilators and angioplasty have been employed.

Competing interests None.
Patient consent Obtained.

REFERENCES