Brainstem haematoma due to presumed cryptic telangiectasia

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SUMMARY Three patients with primary brainstem haematoma are reported. The clinical presentation suggested an initial diagnosis of pontine tumour in two and demyelination in one patient. The subacute course is characteristic of brainstem haematoma due to presumed cryptic telangiectasia, the abnormal vessels being destroyed by the haemorrhage. These findings emphasise the importance of considering haematoma due to cryptic telangiectasia in the differential diagnosis of subacute brainstem lesions.

Cerebral cryptic arteriovenous malformations are telangiectatic lesions, usually measuring less than 3.5 cm in maximum dimensions1 2 and characterised by extremely thin walled capillaries with no muscle or elastic tissue.3-6

The frequency of cryptic malformations of the brainstem is difficult to establish. They are rarely visible on angiography, do not give rise to symptoms or signs unless haemorrhage occurs and the process of haemorrhage itself frequently destroys the abnormal vessels. McCormick4 suggested cryptic telangiectasia within the brainstem are considerably more common than classical arteriovenous malformations, but their frequency has been underestimated by previous necropsy and surgical series.

Three patients with primary brainstem haematoma are now reported. The subacute course emphasises the clinical difficulties in distinguishing haematoma from other causes of subacute brainstem lesions.

Case reports

Case 1
This 27-year-old lady presented in March 1985. Following a 'flu like illness three weeks prior to admission, she noticed difficulty reading "because the words seemed to move". One week later she developed diplopia on right lateral gaze which gradually increased in extent. At the same time she developed persistent nausea and vomiting aggravated by postural change. A week prior to admission she developed paraesthesiae on the right side of the face and within a few hours these spread to involve the trunk and limbs on the left. On the day prior to admission she noted increasing deafness in the right ear with associated tinnitus.

General examination was unremarkable with no cranial

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Fig 1 Case 1 CT scan showing a high density lesion lying in the right side of the brainstem.
bruits. She was fully alert and orientated. The fundi were normal, the right corneal reflex was diminished and there was diminished sensation to touch and pin prick on the right upper lip. There was a right sixth nerve palsy with gaze parietic nystagmus in the left eye on gaze to the right and rotary nystagmus on upgaze. There was a mild right lower motor neuron facial weakness and mild right sensorineural deafness. There was no motor deficit or reflex change. Sensory examination revealed some decrease of light touch and pin prick appreciation on the left sided limbs and left side of the trunk. Vibration and joint position sense was intact. Routine haematology and biochemistry were normal and an initial diagnosis of pontine glioma was made. A high resolution CT scan (fig 1) showed a well defined slightly irregular hyperdense mass lying in the right side of the brainstem. There was a small area of surrounding low density and only slight contrast enhancement. Vertebral angiography showed the basilar artery to be displaced a little anteriorly but no tumour circulation was demonstrated.

Three weeks later a posterior fossa exploration was performed and this disclosed a heaped up discoloured area in the floor of the IV ventricle; the haematoma was incised and evacuated. The histology revealed only a few hyperchromatic cells which were probably reactive in nature. No abnormal vessels were noted. Postoperatively the right facial weakness was worse but in the subsequent four months this has improved considerably. The postural vomiting recovered completely but the remaining clinical signs were unchanged. Examination six months after operation showed partial resolution of the facial weakness.

**CASE 2**

This 27-year-old man presented in June 1984 with a 3 day history of progressively worsening paraesthesiae and numbness in the left limbs and left side of the trunk which came on suddenly following a bout of hicough. Following admission to hospital he developed progressive clumsiness in using the left hand and leg. There had been no visual disturbance or bulbar symptoms. General examination was unremarkable with no cranial bruits. The cranial nerves were normal. There was normal tone and power in the limbs, the tendon reflexes were pathologically brisk but the plantar responses were flexor. There was finger-nose ataxia in the left arm with difficulty in performing fine movements. Sensory examination showed a widening of two point discrimination in the left hand; subjectively the entire left side of the body, except the trigeminal territory, had a sensory disturbance to light touch with cutaneous stimulation causing diffuse dysesthesiae. The gait was normal.

Routine haematological and biochemical investigations were normal and an initial diagnosis of possible demyelination was made. A CT scan (fig 2) showed a circular area of abnormal high density at the level of the medulla, lying slightly to the left. There was slight enhancement. Bilateral vertebral angiography showed lateral stretching of the lateral medullary segments of both posterior inferior cerebellar arteries indicating swelling of the medulla. There was no pathological circulation. A myelogram showed backward displacement of the yallecula confirming the presence of a mass within the medulla.

A posterior fossa craniotomy was performed; the left half of the medulla was swollen by dark, discoloured blood. This was incised and the histology showed only small amounts of necrotic tissue. No abnormal vessels were noted.

One year after operation there has been considerable improvement in the neurological deficit with only minimal dysesthesiae in the left arm and leg.

**CASE 3**

This 22-year-old lady presented in September 1983. Eight days before admission she awoke with a severe headache. On the next day she developed paraesthesiae in the left arm and leg which worsened over the subsequent 48 hours. Over the following three days she developed progressive dysarthria, difficulty looking to the right and weakness of the left arm and leg with some difficulty walking. At the referring hospital a lumbar puncture was performed and within a few hours she became more drowsy and the dysarthria worsened. She was subsequently transferred to the neurological centre.

General examination was unremarkable; there were no cranial bruits. She was very drowsy on admission with dysarthria but was fully orientated. The fields were full, pupils reacted normally and the fundi were normal. The eyes were deviated to the left and did not cross the midline with gaze paretic nystagmus on attempted right gaze. There was diminished pin prick appreciation over all divisions of the right trigeminal nerve and a right lower motor neuron facial weakness. The lower cranial nerves were normal. The tone was increased in the left leg and there was marked poverty of movement of the left limbs which was not formally assessable because of her conscious state. The reflexes were pathol-
There was mild ataxia of gait and limbs with minimal pyramidal weakness on the left. However, there was profound residual posterior column sensory loss in the left arm and leg. Two years following surgery these abnormalities have improved considerably and there is only a mild deficit of joint position appreciation in the left arm.

Discussion

Margolis described four young patients with spontaneous intracerebral haematoma arising from rupture of small vascular malformations. He noted a number of earlier reports of such haemorrhages which had several features in common; in particular they occurred in the younger age group, in the absence of any known predisposing factors and with no demonstrable anatomical cause for the bleeding. Crawford and Russell introduced the term "cryptic arteriovenous and venous haematoma of the brain" to describe small telangiectatic lesions found in 20 patients at necropsy. In a necropsy series of 164 patients with posterior fossa angiomata, McCormick showed that the most common type of malformation in the brainstem was telangiectasia, owing to their relative frequency in the pons. He suggested that cryptic telangiectasia are considerably more common than classical arteriovenous malformations, but their frequency had been previously underestimated by surgical and necropsy studies because the abnormal vessels are usually destroyed by the haemorrhage. Although none of the 27 cases with pontine telangiectasia in his series suffered fatal haemorrhage, he suggested the lesions may account for a proportion of primary pontine haematoma. A number of anecdotal reports have further suggested that the clinical presentation of pontine haematoma due to rupture of cryptic telangiectasia differs from primary pontine haemorrhage.

The three patients described in this series fit the consistent clinical pattern of this condition. All the patients were normotensive and presented in the third decade with the subacute onset of brainstem symptoms and signs over a period varying from three days to three weeks. They suffered hemisensory disturbances and in two patients this was the presenting symptom. Both patients with pontine haematoma developed disturbances of gaze, either as VI nerve or pontine gaze palsy, and facial nerve palsies. An initial diagnosis of pontine glioma was made in two cases and demyelination in one patient. The CT scan appearance of high density lesions in the brainstem suggested the presence of haematoma; in two patients the haematoma impinged on the IV ventricle without rupturing the wall. Angiography showed vessel displacement appropriate to the site of the mass lesion but no abnormalities of vasculature. All the patients...
underwent posterior fossa craniotomy and evacuation of the haematoma although there was clear evidence of progressive deterioration in only one patient. The walls of the haematoma were not biopsied but no abnormal vasculature was found on histological examination of the tissue evacuated. Postoperatively all the patients continued to show moderate neurological deficits, reflecting the preoperative abnormalities, which are progressively resolving.

The natural history of pontine haematoma is uncertain as most reports in the literature concern either post mortem studies or surgical evacuation. Recent reports have indicated that spontaneous pontine haemorrhage may carry a better prognosis than suggested by earlier necropsy reports.15–17 Certainly before CT was available a number of cases of pontine haematoma may have been misdiagnosed as glioma and treated with radiotherapy as in the patient described by Michael et al.24 O'Laire et al.12 have reported successful evacuation in five of six patients and suggested that untreated pontine haematoma is universally fatal. However, a number of series cited by Masiyama14 have reported that the mortality of pontine haemorrhage is between 30% and 68.7% and they found a strong correlation between prognosis and the transverse diameter of the haematoma on CT scan in 26 cases of pontine haemorrhage.

Brainstem haematomas may be misdiagnosed as other causes of brainstem lesions because the clinical course is usually subacute in onset and slowly progressive with occasional remissions and relapses. These diagnoses may include multiple sclerosis, brainstem encephalitis, infarction, granuloma and primary brainstem glioma or metastases.8–13 25. There have been several recent reports of successful surgical evacuation of pontine haematoma.10–12 18–22 The steady progression of symptoms or the exclusion of tumour may be indications for surgical treatment whenever feasible. However, if the clinical state is stable and the cryptic telangiectasia is destroyed there may be a role for conservative management.

Because the abnormal vasculature is frequently destroyed by the haemorrhage, the diagnosis of brainstem haematoma due to cryptic telangiectasia is often circumstantial, based on the subacute clinical course and CT scan findings. The cases reported in this series emphasise the importance of considering haematoma in the differential diagnosis of any subacute brainstem lesion.

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Brainstem haematoma due to presumed cryptic telangiectasia


