Feaver, mucocutaneous haemorrhage, and severe headache during an epidemic of haemorrhagic fever

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A 36-year-old man developed moderate-grade fever, headache, and malaise. On the third day, he noticed purpuric skin lesions and sought medical opinion. Investigations revealed a platelet count of $57 \times 10^9/l$ and haematocrit of 64%. Several people in his locality and the city had similar illness with reported deaths. A diagnosis was made and he was treated with antipyretics for the next 3 days when he developed epistaxis, intense headache, and drowsiness. Cranial computed tomography (CT) (figures 1 and 2) was done and he was referred to this hospital for further management.

On admission he had active epistaxis and complained of persistent headache. History of head trauma in any form was denied. He had no history of any coexisting medical illness. He was febrile, drowsy but oriented, and co-operative. His blood pressure was 130/75 mmHg. He had left-sided hemiparesis (grade 4/5 power) with brisk reflexes. There were no signs of meningeal irritation. Systemic examination did not reveal any abnormality. Laboratory investigations revealed a haemoglobin level of 8.9 g/dl, total leucocyte count was $7.9 \times 10^9/l$ with a normal differential count, haematocrit 57% and platelet count $70 \times 10^9/l$. Blood biochemical parameters and chest X-ray were normal. He was treated with an infusion of platelet-rich plasma, and acetazolamide and mannitol to reduce cerebral oedema. Over the next 4 days, he became afebrile with a rise in platelet count to $130 \times 10^9/l$. However, his neurological status deteriorated with increasing drowsiness, increase in left hemiparesis and bilateral extensor plantar response. A repeat cranial CT scan was not significantly different from the earlier one. In view of his neurological deterioration, he was subjected to a definitive procedure following which he steadily improved. Six months later he was doing well.
The primary disease was dengue haemorrhagic fever (DHF). The World Health Organisation (WHO) definition criteria for DHF during epidemics include fever, muco-cutaneous bleeding, thrombocytopenia, and haemoconcentration. A definitive laboratory diagnosis requires virus isolation or serology. Polymerase-chain-reaction-based diagnosis of the virus has also emerged as a reliable tool for diagnosis, although its routine use has yet to be evaluated.

The pathogenesis of severe complications of dengue virus infections is not clear. Dengue virus infection confers lifetime immunity against re-infection with homologous serotypes, while increasing the susceptibility to other serotypes with a higher risk of complications, such as DHF and dengue shock syndrome (DSS). Studies from Thailand have demonstrated that 90% of all DHF/DSS cases occur with secondary dengue infection. Increased susceptibility to a heterologous serotype is mediated by antibody-dependent enhancement, i.e., sub-neutralising concentrations of antibodies form complexes with the virus and thereby bind to the Fc-gamma-receptor-positive cells such as monocytes through the Fc portion of the immunoglobulin molecule. This enhancement of infection increases the virus load and promotes pathogenesis of DHF/DSS. The secondary antibody response to viral antigens and cytotoxic effect of virus-specific cytotoxic T-lymphocytes sensitised from earlier infection also has an important role in the immunopathogenesis of DHF/DSS. The immune reaction involves activation of the classical complement pathway along with release of various cytokines resulting in increased vascular permeability, disseminated intravascular coagulation (DIC) and antibody-mediated destruction of platelets. These three features are responsible for most manifestations of DHF/DSS.

Common causes of non-traumatic intracranial haemorrhage

- spontaneous intracerebral haemorrhage: hypertension, amyloid angiopathy
- ruptured aneurysm
- ruptured arterio-venous malformation
- bleeding from primary and secondary tumours
- systemic bleeding disorders
- anticoagulation therapy
- drugs: cocaine, amphetamines
- haemorrhagic infarction

Learning points

- hypertension is the commonest cause of all types of spontaneous intracranial haemorrhage
- non-traumatic (spontaneous) subdural haematoma is extremely uncommon
- viral haemorrhagic fevers can be associated with intracranial haemorrhage

Medication. Non-traumatic subdural haematoma has also been reported in association with dural metastasis and tumour embolism. The source of bleeding in two-thirds of cases is arterial. The interval between start of symptoms and diagnosis is variable from 1 day to 5 weeks. The condition has a variable prognosis depending on the size of the haematoma and the intervention provided. Most patients treated in time surgically survive.

Our patient had non-traumatic subdural haematoma and satisfied the WHO criteria for case definition of DHF during epidemics. A definite diagnosis based on serology or viral isolation was not available in this patient, however serologic studies during the epidemic established dengue virus type II as the aetiological agent. The haemorrhagic manifestations of DHF include spontaneous mucocutaneous haemorrhage, namely petechiae, ecchymosis, oozing from venepuncture sites, epistaxis and haematuria. Life-threatening gastrointestinal bleeding occurs in severely ill patients. The pathogenesis of the haemorrhagic manifestations in DHF is immune mediated and involves vascular injury, thrombocytopenia and coagulopathy. Qualitative platelet dysfunction has also been described in DHF. Intracerebral and other forms of intracranial bleeding have not been reported in DHF or, for that matter, in any other type of viral haemorrhagic fever. The pathogenesis of subdural haematoma in the present case is similar to that due to DIC or anticoagulant medication, and represents a hitherto unreported complication of DHF.

Non-traumatic (spontaneous) subdural haematoma as a complication of dengue haemorrhagic fever.
Keywords: subdural haematoma; dengue haemorrhagic fever; haemorrhage


An unusual soft tissue tumour and peripheral eosinophilia

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A 34-year-old woman presented with a painless swelling over her left deltoid region of about 3 weeks duration. There was no history of injury, recent infection or vaccination. The patient occasionally suffered from hay-fever and low-grade joint pain. Clinical examination revealed a firm ill-defined 10 cm x 8 cm non-tender swelling over the posterior aspect of the left deltoid muscle without induration of the overlying skin. No restriction of joint movements was noted.

Initial laboratory studies revealed a normal total white cell count but an elevated eosinophilic count of 12%. Erythrocyte sedimentation rate (ESR) was 19 mm/h. Serum biochemical profile was within normal limits. Rheumatoid factor and antinuclear antibodies were negative. Plain radiograph of the left shoulder only demonstrated soft tissue thickening over the deltoid region. Computed tomography (CT) of the left shoulder is shown in figure 1.

Figure 1 CT scans of the left shoulder, without (A) and with (B) intravenous contrast enhancement

Questions
1 What do the CT scans show?
2 What is the most likely diagnosis?
3 What further investigation is required to confirm the diagnosis?
4 How is the condition managed?
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