Fulminating encephalopathy associated with *Shigella flexneri* infection

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**Summary** Three cases of rapidly fatal encephalopathy associated with *Shigella flexneri* infection are reported. There was a lack of severe intestinal involvement and absence of metabolic derangement. In all 3 patients, areas of necrosis were present throughout the brains; and in one case pontine haemorrhages and demyelination were seen. This report supports the evidence of a particular neurotoxic effect in shigellosis.

Shigellosis is an acute inflammatory disease of the colon produced by bacteria of the genus *Shigella* sp. and characterised by fever, crampy abdominal pain, and frequent loose stools which may contain mucus, pus, or blood. The disease caused by *S. flexneri* is generally benign. Central nervous system complications—such as febrile convulsions—occur in about 30% of patients and may be a reaction to the fever itself, or a manifestation of an acute encephalopathy produced by a neurotoxin elaborated by *Shigella* sp. Here we describe 3 cases in whom infection with *S. flexneri* was associated with a fulminating and rapidly fatal encephalopathy.

**Case reports**

**Case 1.** An 11-year-old boy was admitted to hospital at 0400 hours in a comatose state. He had woken his parents at 0300 hours complaining of abdominal pain and had vomited 3 times. Examination showed blood pressure of 100/60 mmHg, pulse rate 95/min and regular, rectal temperature 37°C, and respiratory rate 12/min with a Cheyne-Stokes pattern. The left pupil was 6 and the right 5 mm in diameter; both were unresponsive to light. There was no papilloedema and no response to caloric or oculovestibular testing. The limbs were flaccid and all reflexes absent. A tip of spleen was palpable and there was a petechial rash on the skin and mucous membranes. *S. flexneri* was subsequently isolated from the stool. Laboratory data were: erythrocyte sedimentation rate 18 mm in the 1st hour (Westergren); haemoglobin 13.9 g/dl; leucocyte count 14.0 × 10⁹/l with normal differential; serum electrolytes and serum enzyme values were normal with the exception of a raised alkaline phosphatase level (189 U/l). The urine showed presence of blood and protein. Lumbar puncture fluid was under slightly raised pressure (200 mmH₂O); cerebrospinal fluid analysis was normal. An electrocardiogram showed a pattern of acute pericarditis (not confirmed at necropsy). Electroencephalogram showed extremely low voltage activity approaching an isoelectric state. Computerised tomography of the brain showed patchy areas of low density, particularly in the temporal lobes. Necropsy showed marked congestion and oedema of the brain, of the small and large bowel, and of the liver, spleen, kidneys, and lungs. The colonic mucosa was congested with a polymorphonuclear cell infiltrate and a fibro- suppurrative exudate containing polymorphonuclear cells covering the mucosal surface. In the brain were found petechial haemorrhages in the pons, patchy areas of necrosis in the grey and white matter, associated with occlusion of small vessels, and areas of perivascular demyelination in the hemispheric white matter.

**Case 2.** A 9-year-old girl was admitted to hospital with a 12-hour history of mild headache, fever, abdominal pains, and the passage of frequent, yellow, loose stools containing neither mucus nor blood. On examination she was pyrexial (38.5°C) with blood pressure 110/60 mmHg and pulse rate 110/min and regular. The stools were at this time loose, yellow, and contained small amounts of blood and mucus. Stool culture subsequently yielded *S. flexneri* type 2a. Laboratory investigations gave normal results. Treatment was started with chloramphenicol 500 mg every 8 hours given orally. Fifteen hours after admission, spontaneous respiration ceased and she failed to respond to noxious stimulation. Pupils were equal (6 mm in diameter) and unresponsive to light; there was no papilloedema. Reflexes were absent throughout the body and the limbs were flaccid. An electroencephalogram performed the next day was isoelectric. Necropsy showed extensive areas of liquefactive necrosis throughout the brain. The rest of the organs were normal apart from the large bowel, the mucosa of which was infiltrated by polymorphonuclear cells.

**Case 3.** An 11-year-old girl was admitted to hospital with a history of sudden onset of headache and fever. Two hours before admission she had vomited 4 times and had passed several loose, yellow stools without blood or mucus. On examination she was drowsy and pyrexial (38°C) with blood pressure...
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SUMMARY Feeding habits before diagnosis were reviewed in 32 infants with salt-losing congenital adrenal hyperplasia who were admitted to hospital in adrenal crisis. Most breast-fed babies failed to thrive, seldom vomited, and despite severe salt wasting, presented at a later age than their formula-fed counterparts.

Salt-losing congenital adrenal hyperplasia (SL-CAH) is a genetically inherited enzyme deficiency, the most commonly affected enzyme being 21-hydroxylase. Clinically, SL-CAH is characterised by a non-specific failure to thrive in the first few days of life that progresses to acute addisonian crisis with anorexia, vomiting, dehydration, dystrophy, diarrhoea, and circulatory insufficiency in the 2nd week. Although the initial crisis occurs in most infants with SL-CAH in the 2nd or 3rd weeks of life, it can occur as early as the first week or as late as several months. Kowarski suggests that the age at which the infant presents depends on the severity of the case, which in turn is related to the degree of sodium