TYPHOID SPINE IN A NIGERIAN WITH SICKLE HAEMOGLOBIN

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Spondylitis has been recognised as a complication of typhoid fever for eighty years but is now becoming increasingly rare in the more developed countries. This paper records a case seen recently in Nigeria. The patient also had sickle cell trait, and the possible significance of this finding is discussed.

CASE REPORT

A Yoruba schoolboy of sixteen was first seen in April 1962, having suffered from fever and pain in the low back and the left hip for three weeks. He was thin but did not look ill. He was able to stand unaided, but his back had lost the normal lumbar curve and was held rigid. Percussion of the lumbo-sacral area caused pain and the erector spinae muscles were tender.

Three weeks later the pain had become very severe and completely prevented him from walking. Radiographs revealed slight diminution of the L.4-L.5 disc space with a suspicion of early bridging (Fig. 1). A diagnosis of acute osteomyelitis of the spine was made. He was admitted and, even though no proven case of staphylococcal infection of the spine had yet been seen at University College Hospital, the staphylococcus at this stage seemed the most likely pathogen, and penicillin and streptomycin were given. To establish a definite diagnosis, a needle was inserted into the L.4-L.5 disc by the right postero-lateral route under radiographic control using an image intensifier coupled to closed circuit television. Very little fluid was aspirated, but salmonella typhi was cultured from the needle and syringe. Typhoid infection was also confirmed by agglutination reactions (Table I), and accordingly chloramphenicol (250 milligrams six hourly) was substituted for penicillin and streptomycin and was continued for six weeks. After five weeks the patient felt much better and suffered only slight back pain. The temperature, which on admission had reached 100 degrees Fahrenheit (37.7 degrees
Centigrade) settled within four weeks. The pulse rate remained consistently between 90 and 100 beats per minute.

**Other investigations**—Haemoglobin electrophoresis showed haemoglobin pattern AS. The Heaf test was 1+. The erythrocyte sedimentation rate was 83 millimetres in the first hour (Westergren). The white blood count two weeks before admission showed: total cells 11,800 per cubic millimetre (polymorphs 47 per cent, eosinophils 33 per cent*, lymphocytes 18 per cent, monocytes 2 per cent). Platelets were plentiful. Sickling was present.

**Progress**—All the symptoms subsided and within three months the spine appeared normal on examination. Nevertheless, serial radiographs showed increasing narrowing of the L.4-L.5 disc space with sclerosis of the bodies and anterior bridging (Figs. 2 and 3).

When questioned further the patient asserted that he had never been ill before, apart from an attack of jaundice shortly before his admission. He had then noticed that his urine was dark and his eyes were yellow for about four weeks. He had received native medicine and had remained at home during this time. In retrospect this was probably an example of one of the many presentations of typhoid fever in West Africans.

The blood continued to agglutinate suspensions of salmonella typhi (Table 1). The white cells never exceeded 11,800 per cubic millimetre and a relative lymphocytosis did not occur.

**TABLE 1**

<table>
<thead>
<tr>
<th>Date</th>
<th>Salmonella typhi D (9, 12)</th>
<th>Salmonella typhi d (H)</th>
<th>Salmonella typhi Vi</th>
<th>Salmonella paratyphi (1, 4, 5, 12)</th>
<th>Salmonella London (3, 10)</th>
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<td>1/2,560</td>
<td>1/160</td>
<td>1/160</td>
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<td>1/1,280</td>
<td>1/320</td>
<td>1/80</td>
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<td>1/1,280</td>
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<td>1/1,280</td>
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The haemoglobin level remained between 11.5 and 14.5 grammes per 100 millilitres. Repeated attempts at blood, stool and urine culture consistently failed to grow either salmonella or any other pathogen. Three months after admission the erythrocyte sedimentation rate had fallen to 11 millimetres in the first hour and cholecystography revealed a normal functioning gall bladder.

**DISCUSSION**

Although Whitman (1930) mentioned a thesis by Maisonneuve on this subject in 1835, the clinical entity of "typhoid spine" was first established by Gibney in 1889. That the specific fevers, notably typhoid fever, could be complicated by osteitis or periostitis had been recorded earlier (Paget 1876, Keen 1877, Terrilllon 1884, Jones 1887).

Osler in 1894 described two cases of typhoid spine but ascribed the pain to a neurosis. Many case reports followed (Lovett and Withington 1900, Cutler 1902, Lord 1902, Favre and Bovier 1913) and Lance (1911) added fifteen new cases to the eighty already reported. Murphy (1916) commented on 164 cases of periostitis and osteitis which occurred in 18,840 patients suffering from typhoid fever (an incidence of 0.82 per cent). Textbooks of orthopaedics

* A common finding in West Africans, and attributed to parasitic infestations.
published during the next fourteen years contain references to typhoid spine (Jones and Lovett 1929, Whitman 1930). Albee in 1919 described the clinical picture and emphasised that the onset is during the early months of convalescence.

During the last thirty years reports of patients with confirmed typhoid spines are rare, and although this may be a result of the widespread appreciation of this condition, it is interesting to note that Huckstep (1962a and b) working with more than a thousand typhoid patients in East Africa only saw one probable case. There are, however, several descriptions of spinal lesions caused by salmonella other than salmonella typhi (Waaler 1935; Anchersen 1947; Rozansky, Ehrenfeld and Matoh 1948; Ralston 1955; Greenspan and Feinberg 1957; Massachusetts General Hospital 1958; Stenström 1958; Waisbren 1960), which often affected the second and third lumbar vertebrae.

Osteomyelitis due to salmonella (either typhoid or non-typhoid), while not very common, is also widely recognised and has been well documented (Ferrier 1952).

Association of salmonella bone infections with abnormal haemoglobins—The presence of sickle haemoglobin, found commonly in many negro races, has in the last few years been shown to bear a strong relation to salmonella bone and joint infections (Hodges, Holt, Jacox and Collins 1951; British Medical Journal 1957; Hook, Campbell, Weens and Cooper 1957; Roberts and Hilburg 1958; Hendrickse and Collard 1960). Patients with sickle cell anaemia (homozygous haemoglobin S disease or SS) are liable to develop infarcts of bone which frequently become infected with salmonella organisms. Similarly, patients with salmonella bone or joint infections are often found to have sickle cell anaemia (SS). A case of "typhoid spine" recently reported from Bamako in West Africa showed the heavy proportion of eighty sickle forms per 100 red cells, and was almost certainly an example of sickle cell anaemia (SS) complicated by salmonella osteomyelitis of the spine (Bourrel and Boissan 1961).

While there is no doubt of the relationship between homozygous haemoglobin S disease (SS) and salmonella bone infections, it is likely that the combination of sickle S and another abnormal haemoglobin C, also found in West Africans and responsible for "sickle cell haemoglobin C" disease (SC), similarly predisposes to salmonella bone infections.

The combination of sickle S and normal adult A haemoglobin producing "sickle cell trait" (AS) has not previously been incriminated as a factor predisposing to salmonella bone infections (Watson-Williams 1962). The association in this patient of sickle cell trait with a typhoid spine is therefore of interest.

TREATMENT

Treatment of typhoid spines is eminently satisfactory, as the condition is largely self-limiting with spontaneous bony fusion of the affected vertebrae. Older writers refer to rest, immobilisation, relief of pain and application of counter irritants, while more recently typhoid osteomyelitis has been found to respond reasonably well to chloramphenicol and surgical intervention (Ferrier 1952). Chloramphenicol alone produced a rapid and dramatic relief from pain in our case. Reports of only two fatalities have been found in the literature; in both of these patients the typhoid spines were complicated by abscess formation and abdominal aneurysm (Miller 1954, Simon and Silver 1957).

SUMMARY

1. A proven case of typhoid spine in a patient with sickle cell trait (AS) is recorded. It responded well to conservative treatment with chloramphenicol.
2. The literature on typhoid spine is briefly reviewed and the relationship between salmonella osteomyelitis and sickle cell disease is discussed.

I wish to thank Dr Hugh Platt for assistance with the interpretation of the laboratory findings, Mr F. Speed for reproducing the radiographs, and Mr H. Jackson Burrows for his valuable advice.

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