SCHISTOMIASIS OF THE SPINAL CORD: REPORT TWO CASES

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SUMMARY

Schistosomiasis affects over 200 million people worldwide. Schistosomiasis of the spinal cord is a rare occurrence. In Africa, there have been recent reports from Egypt and South Africa. In Uganda, the last histological records were over two decades ago. Schistosomiasis of the spinal cord is commonly caused by *Schistosoma mansoni* although *Schistosoma haematobium* has been isolated. Two case reports are presented. In both patients, the diagnosis was made retrospectively. The first patient was a female patient with a lesion in the thoracic region. The second patient was a 21-year-old male with a lesion in the conus. Apart from a block on the myelograms, all other laboratory investigations were negative. The diagnosis was made histologically in both cases with the identification of eggs of schistosoma in the spinal cord. The eggs could, however, not be retrieved from the stool or urine samples. Both patients were treated with antischistosomal drugs and steroids. On follow up they had marked improvement in their neurological signs. We hope to renew attention in this rare but devastating neurological manifestation of a disease which affects many in our region and which if left untreated can lead to permanent neurological damage.

INTRODUCTION

Schistosomiasis affects over 200 million people worldwide (1). Schistosomiasis of the spinal cord is a rare occurrence (2-4). Since 1953 there have been over 40 case reports of schistosomiasis of the spinal cord reported. In Africa, there have been three recent reports from Egypt and five from South Africa. In East and Central Africa, reports have been scanty and were published over a decade ago (6,7).

Schistosomiasis of the spinal cord is commonly caused by *Schistosoma mansoni* (3) although *Schistosoma haematobium* has been isolated (3,5). The pathology of schistosomiasis of the central nervous system is well known (1). Cohen et al (8) found the commonest site in the spinal cord to be the conus medullaris. Early intervention has been found to lead to good neurological recovery (3,9,10).

CASE REPORTS

Two patients who presented to the Neurosurgical Department in 1998 and 1999 are reported. The diagnosis of schistosomiasis of the spinal cord was made retrospectively in both cases.

**Case 1:** Patient A was a young 18-year-old female who was referred from a rural hospital in western Uganda with a history of gradual onset of weakness of the lower limbs, back pain, as well as urinary incontinence developing over 11 weeks. She had a history of dysuria which was being treated as a pelvic inflammatory disease. Clinically she had spastic paraplegia of the lower limbs (Grade 0) with a sensory level at thoracic spine 12 (T12) and localised spinal tenderness at thoracic spine 9 (T9). She was investigated with a full haemogram, thoracolumbar radiographs, CT myelograms and lumbar puncture. The myelogram showed a block at T9 (Figures 1 and 2). The bones appeared normal. At laminectomy a soft intramedullary mass was found. The histology results showed eggs of *Schistosoma haematobium* and myelitis. A urinalysis and stool analysis were performed and the patient started on praziquantel and high dose dexamethasone. Urinalysis and stool results did not yield any ova or cysts. One year later she had normal muscle power and sensation.

**Figure 1**

Plain myelogram of patient A with a block at thoracic spine 9 (T9) (indicated by sudden absence of contrast beneath vertebra pointed at by arrow) The bones appear normal.
The patient was started on praziquantel with high dose dexamethasone and discharged. Follow up at six months revealed recovery of muscle power to Grade 3 and sensory level to T12

DISCUSSION

Patient B was from central Uganda which is not traditionally a schistosomiasis endemic area (11). However, the rest of the presentation was classical (3,9). Patient A was less typical. She had a lesion in the thoracic spine and Schistosoma haematobium was identified as the cause where as most patients are found to have Schistosoma mansoni (2,5). In both patients the diagnosis was not suspected pre-operatively. A study in Natal, South Africa noted that patients who were operated had a clear diagnosis whereas those with myelitis were easily missed (5). Eosinophilia which has been noted in many patients (10,12) was not present in our patients. Peregrino et al (10) have proposed a clinical and laboratory investigation protocol for the diagnosis of schistosomiasis of the spinal cord without a histological diagnosis. This approach may be useful in a patient without mass effects on the spine. Other studies have reported typical lesions on MRI (2,8). However, in a setting with limited facilities, a histological diagnosis of the spinal lesion is often the best option for patients with mass effects and a high index of suspicion should be maintained for patients without mass effects.

The pharmacological treatment of the patients using antischistosomal drugs and steroids is recommended; as well as surgical decompression of tumour-like lesions (9,13,14). However, randomised controlled trials comparing operative and drug treatment have not been performed. No consistent pattern of response to any single treatment modality has been found (15) but it is clear that early treatment leads to subsequent improvement of neurological signs (2,13). We hope that through this report we can renew the interest of physicians in this rare condition which, if not diagnosed, in time can lead to devastating effects.

REFERENCES


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**SOCIETY OF NEUROSCIENTISTS OF AFRICA (SONA) and INTERNATIONAL CHILD NEUROLOGY ASSOCIATION (ICNA) in conjuction with KENYA PAEDIATRIC ASSOCIATION (KPA) and NEUROLOGICAL SOCIETY OF KENYA (NSK) announce**

A Symposium on Child Neurology and Development Medicine in Africa

**Venue:** The Safari Park Hotel, Nairobi, Kenya

**Dates:** 27th – 29th April 2001

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