

23 *Taenia solium* Cysticercosis: Uncommon Manifestations

Gagandeep Singh and Indermohan S. Sawhney

Introduction

The most common manifestations of neurocysticercosis (NC) include seizures, headaches and focal neurological deficits. However, the disorder is known for its pleomorphic presentations. Medical literature is replete with reports of unusual presentations. Knowledge of these uncommon manifestations is important and failure to recognize them often leads to misdiagnosis and delay in management. Several unusual manifestations are reviewed in this chapter with emphasis on spinal cysticercosis.

Spinal cysticercosis

Walton first described the occurrence of a cyst in the ventral portion of the cervical spinal cord at autopsy in 1881¹. Since then, only a limited number of cases of spinal cysticercosis have been described in literature, emphasizing the rarity of this condition. Its incidence at autopsy is about 3%². Several authors consider that autopsy series underestimate the true frequency of spinal cysticercosis on account of the fact that the spinal cord is not routinely examined³⁻⁵. However, spinal cysticercosis is rare in clinical series of cysticercosis as well, its frequency being less than 2.5%^{4,6}. There were

only two cases with spinal involvement in the large series of 450 patients of Dixon and Lipscomb⁶. The condition is likely to be over-represented in neurosurgical series as a number of such cases end up with surgical treatment⁷. Finally, spinal involvement was seen in 5 of 356 cases of a radiological series of NC⁸.

Spinal cysticercosis is classified on the basis of anatomical localization of cysts. The disease may occur at extradural, intradural extramedullary and intramedullary locations. Extradural cysticercosis is extremely rare (Fig. 23.1a and b)⁹⁻¹³. Canelas *et al.* reviewed published literature and collected 35 cases of intradural extramedullary spinal cysticercosis (IDEMSC) and seven cases of intramedullary spinal cysticercosis (IMSC)⁴. Going by their survey of literature, IDEMSC is five times more common than IMSC. Pathologically, the former condition is represented by racemose cysts in the intradural compartment as well as arachnoiditis.

IDEMSC represents the downward migration of intracranial-subarachnoid racemose cysts (Fig. 23.2a-c)^{4,10,14}. Queiroz *et al.* performed elaborate calculations in order to estimate the probability of location of cysticerci in relation to the regional blood supply to various segments of the spinal cord⁵. They concluded that IMSC was acquired through a haematogenous route and

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