A curious case of disseminated cysticercosis in an immunocompetent adult

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Summary  Cysticercosis is an infection with the larval stage of *Taenia Solium* which is estimated to affect over 50 million people worldwide. We report a case of disseminated cysticercosis in an immunocompetent 68-year-old male who presented with back pain, presumed to be musculoskeletal in nature initially. Magnetic-resonance-imaging of the lumbar spine revealed intramuscular (paraspinous and psoas muscles) cysts, innumerable small cystic lesions bilaterally throughout the cerebellar and cerebral hemispheres, midbrain, and right ventricle suggestive of cysticercosis. Treatment with albendazole with dexamethasone for 3 months led to resolution of the cysts with complete resolution of symptoms. Despite its importance, current data on prevalence of this infection, disease burden and the incidence of hospitalization remains incomplete. Mandatory reporting of diagnosis would enable complete understanding of epidemiology of the disease. In this case we have emphasized the importance of early diagnosis of a systemic condition that could have caused serious implications if left untreated.

Keywords: Disseminated, cysticercosis, immunocompetent

Vital signs were normal. On physical exam he had an antalgic gait with tenderness along his left paraspinal muscles with limited extension and flexion of the left leg. The straight leg test was performed and was positive at 20 degrees in addition to decreased strength (4/5) in his hip flexors and quadriceps in the left leg. Complete blood count with differential and basic metabolic panel were within normal limits. Further workup with magnetic resonance imaging (MRI) of the lumbar spine revealed L5 disc protrusion compressing the nerve root in addition to intramuscular (paraspinous and psoas muscles) cysts suggestive of cysticercosis (Figure 1A). MRI of the brain revealed innumerable small cystic lesions bilaterally throughout the cerebellar and cerebral hemispheres, midbrain, and right ventricle consistent with neurocysticercosis in the vesicular stage (Figure 1B). Cysts were additionally seen in the posterior neck musculature and the right temporalis muscle. He was started on albendazole 400 mg twice daily with dexamethasone 6mg twice daily for 3 months which he tolerated well. Repeat MRI of brain and lumbar spine two months after his first MRI showed resolution of the cysts (Figure 1C) with complete resolution of symptoms. The Center for Disease Control and prevention has designated cysticercosis as "one of the five neglected parasite infections". Likewise, the World Health Organization has
designated it as "one of the seventeen neglected tropical diseases" worldwide. A survey was conducted in ED of eleven institutions throughout the US. It was noticed 2.1% of 1,801 patients presenting with seizures were diagnosed with neurocysticercosis. Despite its importance, current data on prevalence of this infection, disease burden and the incidence of hospitalization remains incomplete. Mandatory reporting of diagnosis would enable complete understanding of epidemiology of the disease. In this case we have tried to emphasize the importance of early diagnosis of a systemic condition that could have caused serious implications if left untreated.

DCC is a condition with multiple organ involvement. Although cysticercosis is common, DCC is an uncommon manifestation. It is usually seen in immunocompromised individuals, DCC in immunocompetent state is extremely rare with less than 50 cases being reported in literature, the majority being seen in India. Management of DCC includes albendazole (15 mg/kg/day) for 28 days or praziquantel (10-15 mg/kg/day) for 7-21 days. Further, symptomatic treatment of seizures with antiepileptics and glucocorticoids to decrease the host response and inflammatory changes is recommended.

Our patient was immunocompetent who initially presented with back pain, which was presumed to be musculoskeletal in nature. He was being treated with pain medications along with physical therapy for 4 weeks when he presented again with antalgic gait. At that time spine MRI was done for further evaluation which showed multiple paraspinous and psoas muscle cysts. This emphasize the fact that a high suspicion of disseminated cysticercosis should be considered in immigrant population. When there is suspicion of cysticercosis, a whole body MRI should be done for detection of disease burden and to plan the appropriate therapy accordingly.

References

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(Received August 14, 2019; Revised November 20, 2019; Accepted November 23, 2019)