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Reversible Manifestations of Extraparenchymal Neurocysticercosis

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Reversible manifestations of extraparenchymal neurocysticercosis

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Key Clinical Message

Movement disorders are uncommon manifestations of neurocysticercosis. When present, most are secondary to parenchymal lesions in the basal ganglia. Rarely, movement disorders can occur in racemose/extraparenchymal neurocysticercosis, an aggressive variant frequently associated with cerebrospinal fluid outflow obstruction and hydrocephalus. Appropriate treatment can reverse neurological manifestations.

KEYWORDS

hydrocephalus, movement disorders, neurocysticercosis, neuroinfectious disease, *Taenia solium*

A 21-year-old Ecuadorian woman with previous hydrocephalus due to neurocysticercosis and ventriculoperitoneal shunt placement at age 19 presented with ophthalmoparesis, cerebellar outflow tremor, and bilaterally upgoing toes (Video S1, pretreatment). CSF opening pressure was 16 cm H₂O. Figure 1A shows MRI.

After 3 weeks of treatment with steroids and albendazole, there was clinical and radiological improvement (Video S1, post-treatment; Figure 1B).

Movement disorders are a rare manifestation of neurocysticercosis. Basal ganglia involvement can cause chorea and/or dystonia. Racemose/extraparenchymal neurocysticercosis, an aggressive variant that commonly causes CSF outflow obstruction and hydrocephalus, can present with parkinsonism, cerebellar outflow tremor, cranial neuropathies, and/or corticospinal signs.^{1,2}

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None.

CONFLICT OF INTEREST

None declared.

AUTHOR CONTRIBUTIONS

EMC: involved in the case concept and design, acquisition of data, interpretation of data, and manuscript writing. FDR: performed neurodiagnostic evaluation and critically revised the manuscript for intellectual content. RS: involved in the case concept and design, acquisition of data, interpretation of data, and critically revised the manuscript for intellectual content. DDC: involved in the case concept and design, acquisition of data, interpretation of data, and critically revised the manuscript for intellectual content. JTS: interpreted the data and critically revised the manuscript for intellectual content. MFV: involved in the case concept and design, interpretation of data, manuscript writing, and critically revised the manuscript for intellectual content.

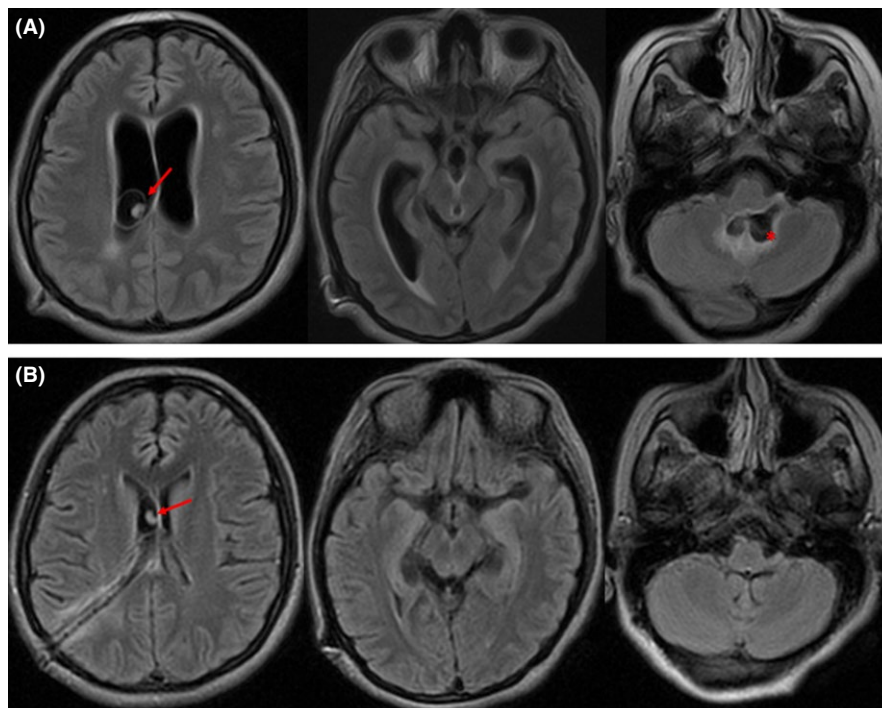


FIGURE 1 MRI FLAIR sequence. A, Pretreatment images demonstrate communicating hydrocephalus with transependymal CSF egress. Note the intraventricular cyst with scolex (arrow) in the right lateral ventricle, and another cyst in the fourth ventricle (asterisk). B, Post-treatment images, obtained 3 weeks later, show resolution of hydrocephalus and fourth ventricle cyst

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SUPPORTING INFORMATION

Additional supporting information may be found online in the Supporting Information section at the end of the article.

REFERENCES

1. Alarcon F, Cedeno Y, de Yebenes JG. Parkinsonism and other movement disorders in 23 cases of neurocysticercosis. *Parkinsonism Relat Disord*. 2017;42:47-53.
2. Garcia HH, Nash TE, Del Brutto OH. Clinical symptoms, diagnosis, and treatment of neurocysticercosis. *Lancet Neurol*. 2014;13:1202-1215.

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