An uncommon cause of seizures in children living in developed countries: neurocysticercosis -a case report

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Abstract
Neurocysticercosis represents an important cause of seizures in children in endemic countries, such as Latin America, Asia and sub-Saharan Africa, while in Europe, especially in Italy, the cases of neurocysticercosis are anecdotal. We report the case of a 6 year old boy, born and lived for four years in Cameroon, who presented a right emiconvulsion. The diagnosis was neurocysticercosis. This case accentuates the need to consider neurocysticercosis in a child presenting with non febrile seizures, mainly if he emigrated from an area of high prevalence or if he had long-term stay in endemic regions.

Background
Neurocysticercosis represents an important cause of seizures in children in endemic countries. It is due to the brain involvement by the larval stage of the cestode Taenia solium (cysticerci). This parasite is commonly found in developing countries of Latin America, Asia and sub-Saharan Africa [1-5]. The prevalence of neurocysticercosis in some of these countries exceeds 10% [6]; conversely in Europe the cases of neurocysticercosis are anecdotal, especially caused by migratory flows from endemic zones or international travels. The case reported below describes a rare cause of seizures in a child who lives in a developed country.

Case presentation
Case report
A 6 year old boy who was born and lived in Cameroon for four years, in a rural area, was admitted to the Emergency Department with seizure lasting more than thirty minutes, not responsive to Diazepam 0.5 mg/Kg e.r. The patient had immigrated to Italy two years previously. No familiarity for seizures or headache. At the age of 1 year the child had experienced an acute and isolated febrile seizure with oculogyration. He had no recent history of traumatic or infective episodes, neither ingestion of drugs or weight loss. When he woke up that morning, he had speaking difficulties, right deviation of the mouth, followed by right head and gaze deviation, right emiconvicusion, and loss of consciousness. At admission the child was afebrile, unconscious, with accelerated heart rate and respiratory difficulties. His head was deviated to the right, there was a generalized hypertonia with hyperextension of right arm and flexion of the left arm. The electroencephalogram (EEG) detected a slow mono-polymorphic activity on the left and central electroencephalographic leads. This activity was absent on the right leads. It suggested a post-critical focal cerebral suffering.

He was directly admitted to the Intensive Care Unit and he was successfully treated with Phenobarbital (5 mg/Kg i.v.). The magnetic resonance imaging (MRI) of the brain showed two cystic round lesions located one in the right lentiform nucleus and one in the left frontal-parietal lobe (Figure 1). In the following 36 hours the EEG displayed an improvement of child’s cerebral conditions with the disappearance of the asymmetric slow activity reported previously. After 48 hours the child was transferred to our Department. Blood exams revealed high eosinophil cell count (720 cells/μL) and Ig E levels (217 UI/ml). Western blot assay detected specific antibodies against cysticercus (LDBIO DIAGNOSTICS, Lyon, France). Therefore the diagnosis of neurocysticercosis was made and the appropriated therapy was started: the child received, orally, albendazole...
(15 mg/Kg/day) in two divided doses and dexamethasone (2 mg/day) for eight days.

Six months later, the eosinophil and IgE levels got back to normal. At brain MRI calcified areas, corresponding to the previous lesions, were indicated, without any new active lesions (Figure 2).

In the following two years the patient remained seizure-free on oral antiepileptic therapy. Then the patient has been lost to follow up.

**Discussion**

Neurocysticercosis is the most common parasitic infection of central nervous system [7]. Taenia solium, responsible of the disease, can causes, in individuals, intestinal infection and cysticercosis. In the classic life cycle humans develop taeniasis after eating not well cooked pork meat, containing cysticerci. Larvae develop into adult tapeworm inside human’s small intestine. The tapeworm releases daily eggs disseminated to the environment through feces. In developing countries, where sanitary regulations are often neglected, roaming pigs may ingest food polluted by eggs, becoming the intermediate host. Once in the pig’s intestinal tract, eggs embrionate into oncospheres, which cross the intestinal wall and, through bloodstream, migrate to the target tissues (brain, muscle, subcutaneous surface). The cycle can be interrupted if individuals accidentally ingest eggs through food or water contaminated by feces. In this case, the parasite perceives the human as an intermediate host and disseminates to a variety of organs, including the central nervous system, where cysticerci develops [8].

The clinical manifestations of neurocysticercosis are variable, depending on the number, stage, location as well as the host response [9,10]. A large part of individuals holding the parasite in the central nervous system are asymptomatic [8]. In some patients symptoms may develop many years after the brain infection, as it has happened to our patient. The most frequent manifestations, due to parenchymal location, are seizures, learning disability, difficulty with balance, behavior changes. Extraparenchymal location of the parasite causes increased intracranial pressure with headache, vomiting and papilloedema [7,8,11].

Parenchymal forms of neurocysticercosis have a better prognosis than extraparenchymal forms, in term of remission of clinical signs [7]. Carpio and Hauser noticed that the risk of seizure recurrence is high after a first acute symptomatic fit but the risk seems related to the persistence of active brain lesions [12].

Histological demonstration of parasites from a brain biopsy should be the safe “gold standard” for the diagnosis of neurocysticercosis. Clearly this procedure is limited because of its invasiveness. Therefore neuroimaging remains the main instrument for the diagnosis. In the identification of extraparenchimal cysts MRI is more sensitive than Computed Tomography (CT) scan [10].

Immunological tests have low sensitivity and specificity, especially for single lesion. The enzyme-linked immunosorbent assay has 65% specificity and 50% sensitivity and it can give false negative results in case of parenchymal neurocysticercosis, inactive lesion or helminthic infections. The enzyme-linked immunoelectrotransfer blot
assay, using purified glycoprotein antigens from Taenia solium cysticerci, has been reported to be highly specific (100%) and nearly 98% sensitive for patients with either multiple active parenchymal cysts or extraparenchymal neurocysticercosis [10,13].

The differential diagnoses include echinococcosis, pyogenic brain abscess, fungal abscess, tuberculoma, toxoplasmosis, a primary or metastatic tumor and infectious vasculitis [14].

Antiparasitic drugs are the mainstay of treatment; in particular, albendazole may be favourable for the treatment of parenchymal cysts because of its power to pass into cerebral spinal fluid [8]. Previous studies recommended the administration of albendazole at a dosage of 15 mg/Kg/die for 1 month, but later studies prove that a 1 week course is equally effective [14]. Corticosteroids are useful for reducing local edema and inflammation around dying parenchymal cysts; so they are often administered together with antiparasitic drugs. Kalra displayed that the association of albendazole and dexamethasone increases complete or partial resolution of lesions and reduces the risk of recurrence of seizure among children presenting one or two ring-enhancing lesion on CT [15].

This case is interesting mainly for epidemiological reasons. In fact neurocysticercosis is endemic in Latin America, Africa and some Asiatic countries. In Europe, many cases have been reported in Portugal, Spain, Poland and Romania. In Italy it is a rare disease. In recent years no cases have been described, but with high rate of immigration from endemic areas (Africa and East Europe) this parasitosis will be found in our country too [16].

Conclusions
This case emphasizes the need to consider a parasitic infection of the central nervous system, particularly...
neurocysticercosis, in a child with an onset of epilepsy especially if he had long-term stay in endemic regions, or he emigrated from an endemic area, although not recently. In case of relevant suspicion of neurocysticercosis it needs testing serum cystercicular antibody and making neuroimaging examinations.

Consent
Verbal informed consent was obtained from the patient’s parents for publication of this case report and any accompanying images, at time of diagnosis; then the patient (with his family) has been lost to follow up.

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Authors’ contributions
IR, CS, FA, SA have made useful contribution in drafting the manuscript and in the revision of the literature. PAT has been involved in revising it critically for important intellectual content. FD has participated in the diagnostic pathways. All authors read and approved the final manuscript.

Competing interests
The authors declare that they have no competing interests.

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