CASE REPORT

Misdiagnosed behavior change revealing fatal primitive intra-cerebral echinococcosis

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Abstract: Central nervous system involvement in hydatidosis is rare compared to other somatic localizations: 1-5%. It is schematically divided into two major types: cerebral involvement and spinal involvement. The brain is a rare localization of hydatid cyst accounting for only 2% of all hydatid disease. The psychiatric manifestations revealing cerebral hydatidosis remain exceptional and often unrecognized and neglected by practitioners despite their potential severity. We report two original observations of fatal cerebral hydatidosis revealed by behavior change (psychomotor agitation, aggressiveness, persecutory delusion, and auditory and visual hallucinations) insufficiently explored in two women aged 45 and 17. The possibility of cerebral hydatidosis must always be evoked in front of any psychiatric symptomatology that remains unexplained in endemic areas for echinococcosis.

Keywords: behavior change, psychiatric manifestations, hydatid cyst, brain hydatidosis, intra-cerebral echinococcosis

1 Introduction

Echinococcosis is the most common cestode in the world with a predilection for the Middle East and North Africa.[1–3] It is still one of the so-called “neglected” tropical diseases[1] and represents a real public health problem as well as a heavy socio-economic burden in several countries of the world.[1, 3, 4] The world’s major endemic regions are the Mediterranean Basin, South America, Asia, including India, Afghanistan, Iran and Turkey; and parts of East Africa.[1–3] Indeed the total number of infested subjects is estimated at 2-3 million worldwide[5] and the severity of this disease lies mainly in its acute complications (rupture, superinfection, anaphylactic shock, etc.) with a risk of sudden death attributed to this parasitosis which remains today “serious and not exceptional”.[6] This mortality mainly affects young people.[5,6]

Brain localizations are part of the so-called unusual or aberrant localizations of this parasitosis (extra-hepatic and extra-pulmonary) observed in less than 10% of cases.[7,8] The psychiatric manifestations revealing cerebral hydatid cyst remain exceptional and often unrecognized and neglected by practitioners despite their potential severity.[9]

We report two original observations of fatal cerebral hydatidosis revealed by behavior change insufficiently explored.

2 Case 1

A woman aged 45 was found dead in her bed in the morning. Her medical history noted that she had been followed for two months for behavioral disorders, family aggression, persecutory delusion and auditory and visual hallucinations of progressive installation. No particular personal or family psychiatric history, drug or substance abuse, or any particular addictive behavior was noted. The somatic examination as well as the basic biological tests requested by her psychiatrist revealed no anomalies. The treatment with neuroleptics prescribed by her psychiatrist was ineffective. Given the unexplained nature of the death, an autopsy was indicated. Postmortem cerebral CT showed a large cerebral mass, of fluid density, intra-parenchymal, supratentorial, occupying the left parieto-occipital lobe, resulting in intracranial hypertension and subfalcine herniation (Figure 1).
The post mortem examination concluded to a broken giant cerebral hydatid cyst. No other anomalies or other associated hydatid localizations were found. The search for toxic substances in the blood and gastric fluid was negative.

Figure 1. Axial cerebral CT: large left parieto-occipital mass of fluid density with intracranial hypertension and subfalcine herniation

3 Case 2

A 17-year-old teenage girl who died suddenly in the emergency room where she was seen for severe acute headaches associated with vomiting.

The interrogation of the family revealed the notion of brutal and unexplained change in her behavior for a month, particularly with psychomotor agitation, aggression towards other family members, and persecutory delusion. She was examined by a psychiatrist and treated with anti-psychotic drugs but without any improvement. No particular family or personal psychiatric history was revealed. Similarly, no recent traumatic event or specific drug or toxic intake has been reported.

Postmortem cerebral CT showed a large multivesicular cerebral mass, of fluid density, intra-parenchymal, supratentorial, occupying the right parieto-occipital lobe and complicated by cranial deformation, intracranial hypertension and a significant mass effect on midline structures (Figure 2).

The autopsy revealed an enormous cerebral hydatid cyst (Figure 3 and Figure 4). No other hydatid localizations were found. The search for toxic substances in the blood and gastric fluid was negative.

4 Discussion

Unusual localizations of human hydatid disease can affect all tissues and organs: the gastrointestinal tract, pancreas, spleen, urogenital system, retroperitoneum, peripheral nervous system, bones, endocrine glands, eyeball, heart, soft tissues ... etc. [7–14]

Central nervous systems involvement in hydatidosis is rare compared to somatic locations: 1-5%. [7–9] It is schematically divided into two major types: cerebral involvement and spinal (spinal cord) involvement. Brain localization was found in 69.2% of central nervous system cases of hydatidosis whereas spinal involvement was found in 30.8%. [15, 16]

The brain is a rare localization of hydatid cyst accounting for only 2% of all hydatid disease [17] and cerebral hydatidosis occupies a significant place among intracranial expansive processes: 3.6%. [18]

Concerning age distribution, children have a significantly higher frequency: from 50 to 93%. [18] This could be explained by the permeability of the ductus arteriosus during the neonatal period allowing the parasite to move from the periphery to the brain. [19] There is no significant difference according to sex, but a slight male predominance is found in a few series. [20] Our two cases are
The hydatid disease can reach any part of the brain but the hemispheres are the preferential localization: 95% of the cases.[21] The cysts are generally located in the supratentorial region in the territory of the middle cerebral artery: the temporo-parieto-occipital region.[22]

The clinical presentation of the hydatid cyst of the brain is similar to that of a brain tumor. It depends on the localization, size and number of cysts. The patient has focal neurological signs (motor deficits such as hemiplegia or hemiparesis, cerebellar syndrome, seizures, disturbances of consciousness) and/or symptoms related to intracranial hypertension (diffuse headaches relieved by jet vomiting and accompanied by visual disturbances).[9, 23] Psychiatric manifestations can be noted in 15 to 20% of cases of cerebral hydatid cyst, particularly at the beginning of their development.[9] These manifestations are nonspecific and are often overlooked and neglected by clinicians.[9, 24, 25]

Several types of behavior disorders have been reported: instability, agitation, violence, aggression, isolation, corporo-sartorial carelessness, visual and auditory hallucinations, schizophrenia-like syndrome, sub-confusion, sleep disorders, hostility, emotional lability, persecutory delusion, hypothalamic stupor, and infantilism.[9, 24, 25]

Hypereosinophilia is rarely found. Similarly, hydatid serology is characterized by low sensitivity in this location; it is rather beneficial for post-operative surveillance to detect recurrences.[16, 18, 23]

The diagnosis of cerebral hydatid cyst by medical imaging (CT and MRI) is generally easy and rarely raises a problem of differential diagnosis since the appearance of cerebral hydatidosis is almost pathognomonic.[21, 26, 27]

The treatment of cerebral hydatid cyst is mainly surgical. However, the use of the anthelmintic agent albendazole is recommended to: sterilize the cyst, reduce the risk of anaphylaxis, reduce the cyst wall stress and decrease the post-operative recurrence.

5 Conclusion

Human hydatid disease of the central nervous system is rare even in endemic areas. The primary brain localization is exceptional and is characterized by a high clinical latency. The psychiatric manifestations associated with these locations are not as rare as thought, but are unknown to clinicians. Our two observations are characterized by the female sex, the late onset in adulthood, the revelation by psychiatric symptoms, and the fatal evolution. The possibility of cerebral hydatidosis must al-
ways be evoked in front of any psychiatric symptomatology that remains unexplained in endemic areas for echinococcosis.

References


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