A seven-year-old boy with a tumour on his right upper leg

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Case

A 7-year-old boy living in a rural area presented with a complaint of swelling in the right upper leg which was present for 1.5 years, but increased in size in the last 3 months and caused pain and difficulty in walking. On physical examination, a soft mass of 4x6 cm without erythema was found in the adductor region of the right thigh. The patient had no history of trauma, surgery or disease.

Eosionophilia (15.4%), leucocytosis (14000/mm³) and a CRP value of 37.7 mg/dL were found in the laboratory tests.

Ultrasoundography (USG) revealed a lesion area with a size of 24x44 mm including extensive internal septations with a thick capsule localized in the muscle in the anterior-medial region of the right thigh compatible with cystic criteria. At this stage, necessary blood tests and magnetic resonance imaging were performed.
Diagnosis: Hydatic cyst

Elisa test performed for echinococcus was found to be positive.

Magnetic resonance (MR) imaging revealed a cystic lesion with regular borders with a size of 28x44 mm including internal septations in the anterior-medial region of the right thigh (Hydatic cyst?). The lesion was hypodense on T1 weighted images, hyperdense on T2 weighted images and showed no marked contrast uptake following IV contrast injection. Contrast uptake was observed in the peripheral region of the lesion (inflammation) (Picture 1).

Postoperatively, 10 mg/kg albendazole was given for 10 days. After obtaining informed consent from the patient operation was performed under general anesthesia and it was observed that the cyst was localized between adductor and sartorius muscles. The cystic mass was excised completely without damaging the mass (Picture 2) and the area was rinsed with hypertonic saline. Afterwards, the mass was opened and it was observed that germinative membrane and hydatid fluid were present (Picture 3). No problem occurred postoperatively. The patient was given 10 mg/kg/day albendazole for three months.

Discussion

Hydatosis is a parasitic infection known since the old times. Four different parasites lead to this disease. These include Echinococcus granulosus, E. Multilocularis, E. Vogeli ve E. oligarthrus. The first two have clinical significance. The other two parasites rarely cause infection in humans (1). The disease is endemic in many regions including Mediterranean countries, the Middleeast, Australia and South America. The annual incidence in Turkey is 12/100000 (2).

Hydatic cyst is found with a rate of 64% in the lungs and with a rate of 28% in the liver in children. In addition, the brain, peritoneal cavity, spleen, heart and bone can also be involved (3). It is believed that primary infection occurring in the muscle tissue in the childhood is due to E. Granulosus. It is observed rarely in children and frequently in the adulthood, because the cyst grows slowly and symptoms arise in later years (4).

Primary skeletal muscle hydatic cyst is observed rarely (with a rate of 0.5-4.7%). The cyst uses oxygen to grow. However, muscles contain lactic acid and this environment is not appropriate for the growth of the cyst (4). The pathogenesis related to skeletal muscle localization of hydatic cyst has not been elucidated fully. Some authors propose that it occurs by bite of an infected dog and some others propose that the embryo reaches the skeletal muscle after passing two organs acting as filters namely the lungs and liver (7).

There are approximately 60 cases of primary skeletal muscle hydatic cyst reported in the literature and the patients are mostly adults (5). It occurs very rarely in children (6).

It is difficult to diagnose primary intermuscular hydatic cyst except for endemic areas. Abscess, hematoma, soft tissue tumor and other lesions should be considered in the differential diagnosis. Biopsy should not be performed, since it would be
harmful in terms of leading live scolexes to enter the circulation and causing anaphylaxis by extending and infecting other compartments. Therefore, primarily imaging is very important in swellings (8).

Our patient is the second pediatric case reported in the literature with its intermuscular localization and a size of 5 cm. Although muscular hydatic cyst is observed rarely, it is still important in urban regions in endemic areas. Radiologic imaging tests and serologic tests may be helpful in the diagnosis of muscular hydatic cyst with an extraordinary localization. It may sometimes be confused with abscess on ultrasonography. Marked hypodense borders of the hydatic cyst can be observed on MRI (9).

Serologic tests can be used to confirm the diagnosis. The common techniques used include indirect immunofluorescence antibody test, ELISA, immunoelectrophoresis and immunoblast tests (10). The diagnosis of our patient was confirmed radiologically, serologically and histopathologically.

Conclusively, hydatic cyst should not be ignored in patients presenting with cystic mass in areas where hydatic cyst occurs endemically and radiologic and serologic tests for hydatic cyst would be performed before performing biopsy. If hydatic cyst is considered, the cyst should be excised totally and drainage should not be done.

References