Hydatid cyst disease caused by Echinococcus granulosus may at times affect unusual sites and present a diagnostic challenge especially if found at rare locations. A 51 year male presented with a slowly progressive swelling in the upper medial aspect of the left thigh without pain and any discharge. There was no history of trauma and any dog keeping. Examination revealed a large 10 × 12 cm soft tissue swelling which was non-tender, non-pulsatile, without any features of acute inflammation figure 1. Overlying skin was normal. No bruit or venous hum was heard.

Routine blood investigations, ultrasound of the abdomen and X-ray of chest was normal. Ultrasound of the area showed multicystic swelling. FNAC of swelling revealed a cystic lesion and reported as parasitic or lymphatic cyst. On serological tests, IgG for hydatid was raised with a value of 74.20 IU. MRI of thigh revealed multiple, large, multicystic well defined masses in adductor muscle compartment in the left thigh. Largest cyst was 83 × 69 × 103 mm in size. Multiple, variable cysts were seen divided by septa. No obvious bone or joint involvement was noticed figure 2. MRI features were suggestive of hydatid mass.

Patient was operated under general anesthesia. Hydatid cyst of approximately 25 × 15 × 15 cm was found to be located in the medial aspect of left thigh with multiple daughter cysts in the adductor muscle compartment figure 3 and 4. No neurovascular involvement was found. Care was taken to avoid spillage. 10% Betadine soaked mops were placed around the cyst and hypertonic saline (3% NaCl) instilled in it. Pericystectomy with drain placement done.

Post-operative course was uneventful. Drain was removed on post-operative day four. Patient was discharged on antihelminthic Albendazole 400 mg twice a day for 1 month. Histopathology report revealed a germinal layer, lamellated ectocyst with fibrous outer layer. Marked foreign body giant cell reaction was observed, confirming the diagnosis of hydatid cyst.

Primary hydrated cysts in muscle are an uncommon manifestation, accounting for only 3% of all patients with hydatidosis [1]. Muscular hydatidosis usually are found to occur as isolated lesions without hepatic or pulmonary lesion [2]. The most common locations for development of parasitic cysts are the liver (65%-75%) and lung (25%-30%) [3], the involvement of other organs including brain, heart, kidney, bone, skeletal muscle, breast, thyroid comprise only about 10% and are considered as unusual presentation [4].

Growth of cyst in different organs of body is variable and is dependent on patient factors, parasite factors, host reaction and presence of any complications. Compressible organs like lung and brain facilitate the growth of cyst [5]. Constant contraction of muscles and production of lactic acid probably retards the growth of the cyst in the muscle. Hydatid
cysts in liver and lung. This represents the parasitic membrane (germinal layer) and a collagen rich membrane as a host response (pericyst). Comert et al. first described the “water lily sign” as a pathognomonic sign for intramuscular hydatid cyst [10], the collapsed parasitic membranes which are hypointense on all T1 weighted images represent a non viable cyst and is described as a “Snake sign or Serpent sign” [11].

The treatment of muscle hydatid cyst is surgical excision i.e. pericystectomy while avoiding spillage of hydatid fluid and daughter cysts. This should be followed by antihelminthic medication to prevent recurrences. Mohamed et al. have analysed the efficacy of albendazole alone versus combined albendazole and praziquantel in human hydatidosis are of the view that praziquantel prevents encystment of protoscolecies following perioperative spillage [12]. Percutaneous aspiration, infusion of scolicidal agent and reaspiration (PAIR) under imaging is another option available for inoperable cases.

Adductor muscle hydatidosis is a rare presentation of this disease and has to be included in the differential diagnosis of soft tissue swellings.

References