Cysticercosis of foot: A rare diagnosis in a rare location

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Abstract
Cysticercosis is an infection caused by the larval form of tape worm Taenia Solium. This infection commonly presents with central nervous system involvement. Here we present a case of isolated cysticercosis of foot diagnosed on Magnetic Resonance Imaging (MRI). The case was successfully managed conservatively.

Keywords: cysticercosis, tapeworm, foot, albendazole, MRI

1. Introduction
Cysticercosis infection in humans is caused by Taenia Solium[1]. The cysts occur first in central nervous system followed by the eye, striated muscles, subcutaneous tissue and rarely in other locations. To our knowledge presence of isolated cysticercosis infection in rare[3-6]. Now we present this case of isolated cysticercosis of foot which was diagnosed accurately on Magnetic Resonance Imaging (MRI) and resolved successfully on treatment with albendazole.

All the blood counts were normal. Erythrocyte sedimentation rate was elevated 75mm/hr. No other positive laboratory reports were present.

MRI foot showed small cystic scolex shaped lesion measuring 8mm in the plantar aspect deep to flexor tendons at the level of 2nd metatarsal with a hypo tense focus and associated collection, possibility of soft tissue cysticercosis.

Computed tomography (CT SCAN) of brain of has been done to rule out any cerebral lesions. Report showed normal brain study.

Patient was managed conservatively with anti-helmintic drug Albendazole 15mg/kg/day in two divided doses for three weeks, diclofenac as an analgesic. She was given one dose of dexamethasone 4mg injection intra muscularly. Patient was followed on every third day for three weeks. Pain and edema of foot subsided in a week. Swelling gradually reduced in three weeks’ time. After one and half month patient was completely relieved of symptoms.
Figure 1: Clinical photographs showing edema of foot and cystic swelling

Figure 2: Oblique and anterio-posterior radiographs of foot

Figure 3: MRI Images both sagittal and coronal sections showing scolex shaped cystic lesions suggestive of soft tissue cystercerosis with associated abscess
3. Discussion

Cystercerosis infection in humans is caused by *Taenia Solium*[1]. It is common in Asians (mostly Indians) and African population where the pigs are raised for consuming purpose and socio economic status is low[7].

In the life cycle of tape worm humans are the definitive host for tape worm infection and the tape worm enters humans generally through inadequately pork or beef consumption. Ingested eggs hatch in the small intestine, releasing oncospheres that penetrate the bowel mucosa and enter the bloodstream to travel to various tissues where they develop to form an encysted larval form of *T. Solium* known as cystercerosis cellulosae. When the larva dies, it induces an aggressive granulomatous inflammatory response, leading to characteristic organ-specific symptoms.

The cysts occur first in central nervous system followed by the eye, striated muscles, subcutaneous tissue and rarely in other locations. Presence of multiple muscular cysts and isolated involvement in foot is rare. So this case can be considered as a rare disease at a rare location.

Mostly this type of cystercerosis infection in foot is a rare in possibility. This is the first case in 25 years from the department of orthopaedics existence in R.L Jalappa Hospital, Tamaka, Kolar.

This case mostly goes unnoticed and will be asymptomatic. The swelling in due to release of antigens after the death of disease. In our case any trivial trauma to the foot in the region of cyst which was unnoticed might have caused the release of antigens resulting in death of inflammation.

Three different clinical manifestations of muscular cystercerosis are described: myalgic myopathic type; the nodular or mass like type; and the pseudohypertrrophy type in which multilocular cyst formation occurs in group of muscle[8][9]. The myalgic type results from death of the cyst and leakage of fluid leading to inflammation. The nodular type or pseudotumor type both result from degeneration of the cyst and slow intermittent leakage of fluid over time, leading to a chronic inflammatory response with collection of fluid around the cyst producing a mass. This case is a myalgic variant.

Del Brutto et al proposed diagnostic criteria for human cystercerosis[10]. They suggested that apart from Immunochemical studies (including detection of anticysterceral antibodies), CT and MRI are important tools in diagnosis. Although MRI is more specific for neurocystercerosis, CT may be the modality of choice for muscle cysts as it can demonstrate multiple cysts in a honeycomb or leopard skin pattern against a background of muscle mass.

Treatment of cystercerosis depends on the site involvement. It is generally treated with antiparasitic drugs in combination with inflammatory drugs[11]. Surgery is sometimes necessary to treat cystsin certain locations when patient is not response to drug treatment. Even if you don’t treatment to treat parasite, you may need treatment for the symptoms caused by the infection, such as medication to reduce of seizures you have.

Albendazole acts by inhibiting microtubule formation. The loss of cytoplasmic microtubules blocks glucose uptake in the larval and adult stages of the parasites, thereby depleting their energy reserves and causing death; So treatment with albendazole along with anti inflammatory successfully helped in resolving the disease.

4. Conclusion

So here Cystercerosis of foot is a rare diagnosis in a rare location which was successfully treated non operatively with use of albendazole and anti inflammatory drugs.

References


