

E-mail :info@kistmcth.edu.np I www.kistmcth.edu.np
Journal of KIST Medical College

# Cysticercosis of gastrocnemius muscle

## Rajesh Bahadur Lakhey<sup>1</sup>, Sharad Chandra Adhikary<sup>1</sup>, Dinesh Kafle<sup>1</sup>

<sup>1</sup>Department of Orthopedics, Tribhuvan University Teaching Hospital, Institute of Medicine, Maharajgunj, Kathmandu, Nepal

## ABSTRACT

Cysticercosis is a common parasitic infection in our part of the world involving the central nervous system, adnexal structures of the eye, skeletal muscle, and subcutaneous tissue. The principal mechanism of transmission is through ingestion of Taenia Solium eggs or contamination of fruits and vegetables fertilized with contaminated faecal materials. Solitary intramuscular cysticercosis, without symptoms of central nervous system involvement is rare. We present a case of solitary intramuscular cysticercosis involving gastrocnemius muscle in a 12-year old boy, a rare picture without any neurologic or systemic manifestation.

Keywords: Cysticercosis, Gastrocnemius

Citation: Lakhey RB, Adhikary SC, Kafle D.Cysticercosis of gastrocnemius muscle.JKISTMC 2021;3(1)5:33-35

## **INTRODUCTION**

Cysticercosis is caused by encysted larvae of tapeworm Taenia solium, the pork tapeworm. This disease is endemic in our part of the world. Mostly central nervous system (60-90% of the cases) is involved.1 It is rare to have isolated muscular involvement without the symptoms of central nervous system involvement , and may cause diagnostic dilemma. Our patient presented with rarely seen isolated intramuscular mass in gastrocnemius muscle

### **CASE REPORT**

A 12-year old child presented with mass in left calf for 1 year which rapidly increased in size and became painful for one month . There was no history of fever and trauma. For the investigation, Ultrasound and MRI was done . Ultrasound showed cystic mass with a hyperechoic shadow, suggesting a parasitic cyst with scolex.(Figure 1) MRI showed relatively well- defined T1 low and T 2 high signal intensity lesion , with T1 hyperintense rim with surrounding muscular and tendinous edema , which were the features of myocysticercosis with infection, or muscular

#### Correspondence:

Rajesh Bahadur Lakhey, Department of Orthopedics, Tribhuvan University Teaching Hospital, Institute of Medicine, Kathmandu, Nepal Email: <u>neprajesh@hotmail.com</u> Conflict of interest: None Source of support: None **Article info** Received: 17 Jan 2021 Accepted: 25 Jan 2021 Published: 31 Jan 2021

#### Copyright

JKISTMC applies the Creative Commons Attribution- Non Commercial 4.0 International License (CC BY) to all works we publish. Under the CC BY license, authors retain ownership of the copyright for their article, but authors allow anyone to download, reuse, reprint, distribute, and/or copy articles in JKISTMC, so long as the original authors and source are cited.



#### JKISTMC Jan 2021 Vol 3 No 1 Issue 5:33-35

strain with surrounding muscular edema. (Figure 2) FNAC was done, which showed acute inflammatory cells suggesting infection . Excisional biopsy was planned . During surgery , a woody mass was found in the gastrocnemius muscle , which was excised . On cross section , liquefaction of the center of the mass was seen. On histopathological examination , the features of myositis , with eosinophil rich inflammation and vague granuloma , which suggested parasitic lesion was found. The patient was prescribed 4 weeks of Albendazole and he improved uneventfully.

#### DISCUSSION

Human cysticercosis is a tapeworm infection caused by T. solium. This condition is common in our part of the world .2 It is considered the most common parasitic disease of the central nervous system in immunocompetent individuals .<sup>3,4</sup> Less frequently, other organ systems can also be involved, including skeletal muscle, subcutaneous tissue, the eyes, the tongue, the oral cavity, the breast, the heart, and the lungs.<sup>5,6,7</sup> In the normal life cycle of T. solium, people are definitive hosts and pigs are intermediate hosts. Sometimes, human beings also become the intermediate hosts by ingesting eggs from contaminated food or water <sup>3,8,9</sup> Cysticerci form in human beings, in central nervous system and less commonly in the eyes, lungs , heart , oral cavity , or breast .5,6,7 The encysted larvae may remain viable for

years and usually produce no symptoms.

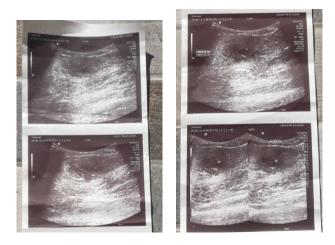


Figure 1. Ultrasound of the lesion



Figure 2. MRI of the lesion

Ultimately , the larvae die; this induces a granulomatous inflammatory response that may produce symptoms, depending on the anatomic site.4, 5,10 Isolated muscular

involvement is rare in cysticercosis.<sup>11</sup> There are very few cases with isolated muscular cysticercosis reported in the literature. <sup>4,10,12–16</sup>

The intramuscular cysticercosis may present clinically as myalgia, pseudotumor, soft tissue tumour, lipomas, epidermoid cysts, granular cell tumours, neuroma, neurofibromas, sarcoma, myxoma and pyomyositis.<sup>17</sup>

Three different clinical manifestations of muscular cysticercosis are described: myalgic myopathic type; the nodular or mass like type; and the pseudohypertrophy type in which multilocular cyst

formation occurs in group of muscle. 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.<sup>18</sup>Our case was characteristic of the myalgic variant.

#### JKISTMC Jan 2021 Vol 3 No 1 Issue 5:33-35

#### REFERENCES

- Ramraje S, Bhatia V, Goel A. Solitary intramuscular cysticercosis – a report of two cases. Australas Med J. 2011;4(1):58–60
- Singal R, Mittal A, Gupta S, Gupta R, Sahu P, Gupta A. Intramuscular cysticercosis diagnosed on ultrasonography in thigh: A rare case report. North Am J Med Sci 2010; 2:162-4.
- Del Brutto OA, Sotelo J. Neurocysticercosis: an update. Rev Infect Dis. 1988;10:1075–1087.
- Ogilvie CM, Kasten P, Rovinsky D, et al. Cysticercosis of the triceps: an usual pseudotumor. Case report and review. Clin Orthop. 2001;1: 217–221.
- Brown WJ, Voge M. Cysticercosis: a modern day plague. Pediatr Clin North Am. 1985;32:953–969.
- Malik SR, Gupta AK, Choudry S. Ocular cysticercosis. Am J Ophthalmol. 1968;66:1168– 1171.
- Romero de Leon E, Aguirre A. Oral cysticercosis. Oral Surg Oral Med Oral Pathol Oral Radiol Endod. 1995;79:572–577.
- Yamashita P, Kelsey J, Henderson SO. Subcutaneous cysticercosis. J Emerg Med. 1998;16:583–586.
- Botero D, Tonawitz HB, Weiss LM, et al. Taeniasis and cysticercosis. Infect Dis Clin North Am. 1993;7:683–697..
- Kazanjian PH, Mattia AR. Case 26-1994: a 20-year-old Philippine woman with soft tissue mass in the forearm. N Engl J Med. 1994; 330:1887–1893.
- 11. Zemeno-Alanis GH. A classification of human cysticercosis. In: Fissler A, Willms K, Laclette JP, et al, eds. Cysticercosis: Present State of Knowledge and Perspectives. New York: Academic Press; 1982:107–127.

- Anderson GA, Chandi SM. Cysticercosis of the flexor digitorum profundus muscle producing flexion deformity of the fingers. J Hand Surg [Br]. 1993;18:360–362.
   Kung IT, Lee D, Yu HC. Soft tissue cysticercosis. Diagnosis by fineneedle aspiration. Am J Clin Pathol. 1989;92:834–835.
- Brown ST, Brown AE, Flippa DC, et al. Extraneural cysticercosis presenting as a tumor in a seronegative patient. Clin Infect Dis. 1992;14:53–55.
- Yue X. Fine-needle aspiration biopsy diagnosis of cysticercosis: a case report. Acta Cytol. 1993;38:90–92.
- Abdelwahab IF, Klein MJ, Hermann G, et al. Solitary cysticercosis of the biceps brachii in a vegetarian: a rare and unusual pseudotumor. Skeletal Radiol. 2003;32:424–428.
- Rangdal SS, Prabhakar S, Dhatt S S, Prakash M, Dhillon MS. Isolated Muscular cysticercosis: A Rare Pseudotumor and Diagnostic Challenge, can It be treated Nonoperatively? A Report of Two Cases and Review of Literature. J Postgrad Med Edu Res 2012; 46 (1): 43-48.
- 17. Horton J. Biology of tapeworm disease. Lancet. 1996; 348:481. .
- Mittal A, Das D, Iyer N, Nagaraj J and Gupta M. Masseter cysticercosis- a rare case diagnosed on ultrasound. Dentomaxillofac Radiol. 2008; 37:113-6.