

Subcutaneous Cysticercosis Mimicking Soft Tissue Neoplasm – A Case Report

Lalawmpuii Pachuau¹, Chawngthu Vanlalhlua²

¹Senior Pathologist, Department of Pathology, Civil Hospital, Aizawl, Mizoram, India.

²Associate Professor, Department of Surgery, Zoram Medical College, Mizoram, India.

Corresponding Author: Dr Chawngthu Vanlalhlua, M.S, FAIS, FMAS, EFIAGES, FIAGES, FDFM

Abstract

Introduction: Cysticercosis is an infection of both humans and pigs with the larval stages of the *Taenia solium*. This infection is caused by ingestion of eggs or gravid proglottids shed in the feces of a human tapeworm carrier. The subcutaneous form of the disease is a relatively rare clinical entity. **Aim:** The aim of this paper is to report a case of Subcutaneous cysticercosis involving the left anterior chest wall mimicking soft tissue neoplasm. **Material and Method:** This is a case of 35 years old female with complaints of lump in left anterior axillary line associated with itching for the last 5 months. The lump slowly increases in size with mild pain. On clinical examination, the lump was located subcutaneously over left lateral border of pectoralis major and firm to hard in consistency with mild tenderness. No skin change noted. Fine needle aspiration cytology was signed off as suspicious of Histiocytic tumour. Excision of lump was done under local anaesthesia and sent for histopathological examination. Histopathological examination revealed cysticercus cellulose parasite with an extensive mixed inflammatory cell infiltrate in the surrounding tissue. The patient was prescribed oral antihelminthic therapy. **Conclusion:** Subcutaneous cysticercosis is a relatively rare form of cysticercosis but should always be borne in mind during the evaluation of subcutaneous swellings. In any lumps with a subcutaneous cyst showing central cystic area containing a caseating- like whitish fluid with covering wall, histopathological examination should always be done to confirm the diagnosis.

Keywords: Subcutaneous Cysticercosis, *Taenia solium*, Histopathological examination

Date of Submission: 18-07-2020

Date of Acceptance: 02-08-2020

I. Introduction

Cysticercosis is an infection of both humans and pigs with the larval stages of the *Taenia solium*. This infection is caused by ingestion of eggs or gravid proglottids shed in the feces of a human tapeworm carrier¹. Though it is endemic in virtually all developing countries in Central and South America, Asia, and Africa, the subcutaneous form of the disease is a relatively rare clinical entity. Humans are infected either by ingestion of food contaminated with feces, consumption of infected raw vegetables, contaminated water, consumption of undercooked pork that harbors the larva, autoinfection and regurgitation of gravid proglottids^{1,2}. Once eggs are ingested, oncospheres hatch in the small intestine, invade the intestinal wall and migrate to striated muscles, brain, liver and other tissues where they developed into cysticerci¹. It has a worldwide distribution and is potentially harmful with variable clinical manifestations depending the location. It can be diagnosed on serology and radiology but confirmation is on histopathological examination of the involved tissue biopsy specimen³.

II. Case Presentation

We presented a case of 35 years old female with complaints of lump in left anterior axillary line associated with itching since January 2020. The lump slowly increases in size and so patient went to a hospital where fine needle aspiration cytology was performed. The report was signed off as suspicious of Histiocytic tumour. On clinical examination, the lump was located subcutaneously over left pectoralis major and firm to hard in consistency with mild tenderness, overlying skin was intact, there was no redness. Further examination showed no other lesion at other sites. Excision of lump was done under local anaesthesia maintaining a strict aseptic measures. Patient was sent home with oral antibiotics and painkiller prescription. The excised lump was firm, greyish white measuring 3x2x4 cm. On serial slicing, cut surface shows central cystic area containing a caseating like whitish fluid. Hematoxylin-eosin stained sections show a cyst wall lined by eosinophilic material that is thrown into folds. Beneath this is myxoid degeneration of cyst wall, occasional eosinophilic spherules was also present. Surrounding stroma shows dense inflammatory infiltrates composed predominantly of foamy histiocytes which could have misguided the cytologist into believing the lesion to be of a benign histiocytic

tumour, the other components are lymphocytes, eosinophils, polymorphs and plasma cells. After the histopathological report came out, patient was prescribed oral antihelminthic therapy.



Fig 1: Post operative patient on follow up



Fig 2: Excised specimen of cyst wall



Fig 3: Excised specimen of cyst wall

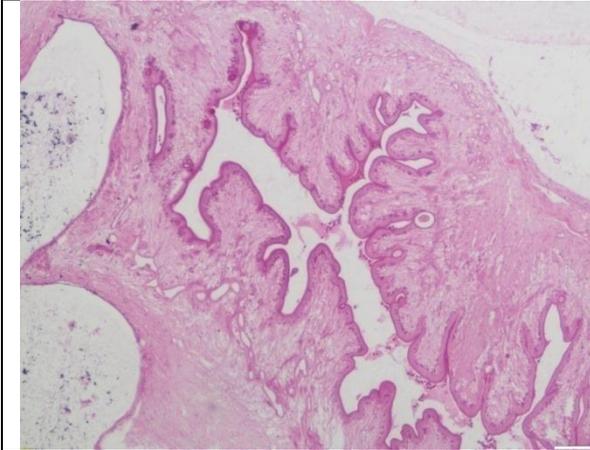


Fig 4: Cyst wall lined by eosinophilic material that is thrown into folds. H&E, x 5

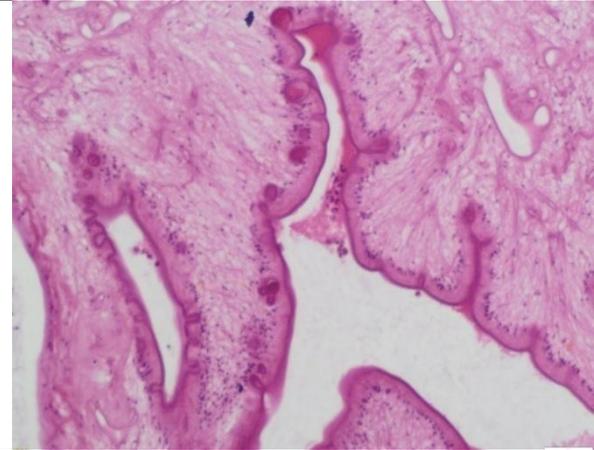


Fig 5: Lining eosinophilic material on high power. H&E, x10

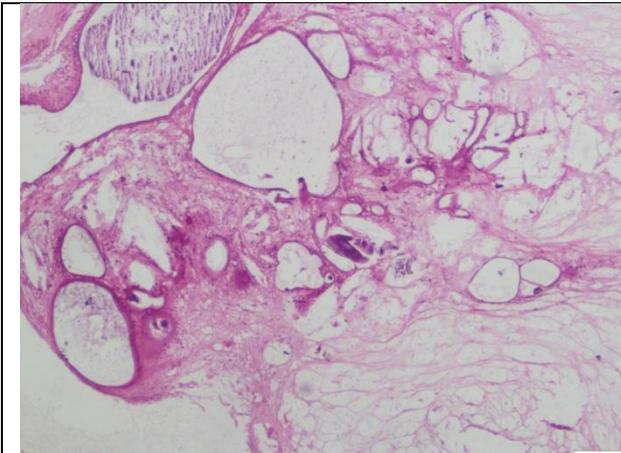


Fig 6: Myxoid degeneration of cyst wall. H&E, 10x

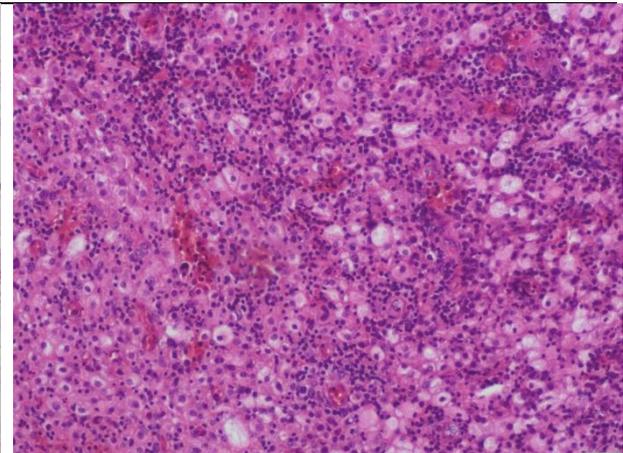


Fig 7: Dense inflammatory infiltrates composed of foamy histiocytes, lymphocytes, plasma cells eosinophils. H&E, 10x

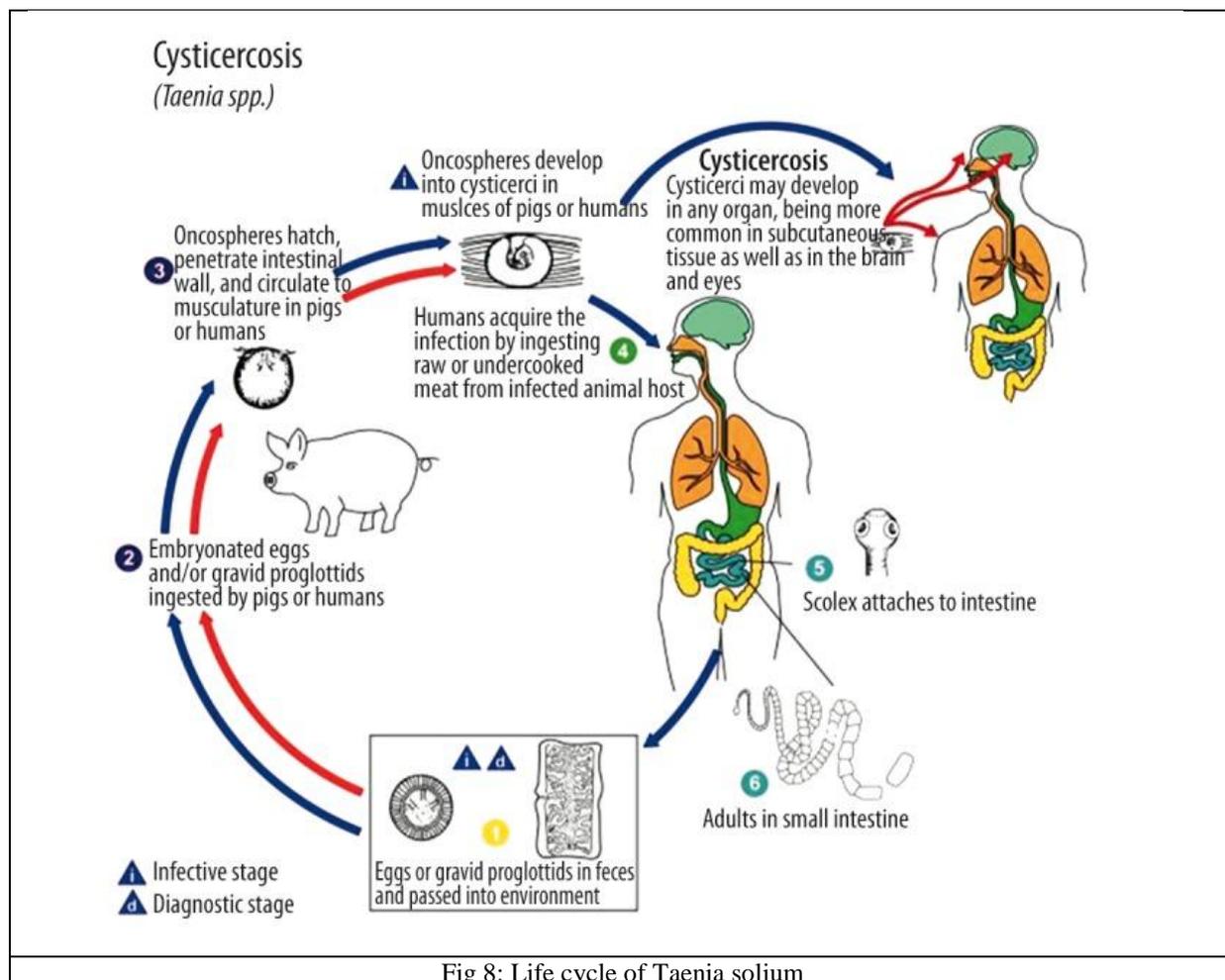


Fig 8: Life cycle of *Taenia solium*

III. Discussion:

In case of infection with *Taenia solium*, humans act as both the obligate definitive host and intermediate host. Infections as a definitive hosts rarely cause significant clinical problems. Whereas, infections in humans as intermediate hosts often causes serious clinical sequelae or death⁴. Cysticercosis is a parasitic infection caused by the larval stages of the pork tapeworm, *Taenia solium*^{5,6}. Human cysticercosis has a worldwide distribution including Central and Eastern Europe, South America, Africa and tropical countries like India⁷. The perpetuation of this parasitic disease is related to poor sanitation and hygiene^{8,9}.

Humans can become infected by consuming gravid proglottids either through feco-oral route or by autoinfection^{10,11}. There is also a high risk of infection by reverse peristalsis resulting in internal regurgitation of the eggs into the stomach when the intestine harbours a gravid worm^{12,13}. The oncospheres penetrate the intestinal mucosa and develop into cysticerci after getting carried to various parts of the body including brain, eyes, striated muscles, liver, heart, lungs, peritoneum, breast and subcutaneous tissues. Ingestion of the undercooked pork containing these cysticerci is the exclusive path to the development of human intestinal *Taenia solium* tapeworms. By means of a single scolex or head, these cysts attach to the small intestine. Adult tapeworms develop and reside in the small intestine for a period varying from months to years. Basically, all the clinical symptoms can be attributed to the vigorous granulomatous inflammatory reaction that occurs when the larvae die¹⁴.

Cysticercosis is commonly seen in the brain and eyes, which together constitute 86% of these cases^{15,16}. The remainder is mainly located in the muscles, heart, lungs, peritoneum and breast^{17,18}. Subcutaneous cysticercosis is a relatively rare form of cysticercosis but should always be born in mind during the evaluation of subcutaneous swellings. It can be confused with other clinical entities depending upon the location of the swelling like a lipoma, ganglion cyst, sebaceous cyst, dermoid, abscess, pyomyositis, tuberculous lymphadenitis, neuroma, sarcoma, myxoma, neurofibroma or fat necrosis¹⁹.

The clinical features of subcutaneous cysticercosis depend on the location of the cyst, the cyst burden, and the host reaction^{20,21}. It may cause painless or painful subcutaneous nodules¹². Lymphadenopathy is a rare

mode of presentation of cysticercus infestation¹⁷. Lohra S. et al. first documented a case of subcutaneous cysticercosis involving the left anterior chest wall with ultrasound evaluation²².

Serological tests for detecting antibodies against cysticercosis are used to confirm the diagnosis. Enzyme-linked immunoblot assay is more sensitive and specific than ELISA (Enzyme linked immune sorbent assay)²³. However, sensitivity of serological tests tends to be high for patients with multiple cysts (94%), but substantially lower for patients with a single cyst or calcified cysts (28%)²⁴.

The diagnosis of cysticercosis can be confirmed by fine-needle aspiration cytology (FNAC) or biopsy, which shows the detached hooklets, scolex, and fragments of the spiral wall of cysticercosis cellulosa²⁵. Biopsy and histologic examination containing the cysticerci is the most definitive method of diagnosis²⁶.

Treatment of subcutaneous cysticercosis depends on the location of the cysts²⁷. Surgical excision is done for isolated soft tissue cysticercosis associated with an abscess²⁸. Cysts that are not associated with an abscess can be treated with antihelminthic medications such as albendazole or praziquantel²⁹.

IV. Conclusion:

Subcutaneous cysticercosis is a relatively rare form of cysticercosis but should always be borne in mind during the evaluation of subcutaneous swellings. Human cysticercosis has a worldwide distribution including Central and Eastern Europe, South America, Africa and tropical countries like India. Parasitic infections should be considered in the differential diagnosis of soft tissue lesions, especially in endemic populations. Health education and good sanitary practices helped in preventing transmission of infection. In India February 10th is observed as National Deworming Day which is an initiative of Ministry of Health and Family Welfare, with a mop up day on February 15th. In any lump with a subcutaneous cyst showing central cystic area containing a caseating - like whitetish fluid with covering wall, histopathological examination should always be done to confirm the diagnosis.

References:

- [1]. Lynne SG: Parasitology case history 5 (histology). Available from <http://www.med-chem.com>. Accessed July 19, 2020.
- [2]. Rajwanshi A, Radhika S, Das A, Jayaram DN, Banerjee CK. Fine-needle aspiration cytology in the diagnosis of cysticercosis presenting as palpable nodules. *Diag Cytopath* 1991; 2(14); 7(5) 517-519.
- [3]. Sawhney M and Agarwal S: Cysticercosis: hooked by a hooklet on fine needle aspiration cytology-a case report. *Case Reports in Infectious Diseases*. Vol 2013, Article ID 315834, 2 pages
- [4]. Lightowlers MW. Cysticercosis and Echinococcosis. *Current Topics in Microbio and Immuno* 2012; 365: 315–335.
- [5]. Sawhney M, Agarwal S. Cysticercosis: Hooked by a hooklet on fine needle aspiration cytology – a case report. *Case Rep Infect Dis*. 2013;2013:315834.
- [6]. Ghimire PG, Ghimire P, Rana R. Spectrum of typical and atypical clinico-histopathological and radiological presentation of soft tissue and muscular cysticercosis in Mid-Western and Far-Western Region of Nepal. *J Clin Diagn Res*. 2015;9(9):EC01–3.
- [7]. Neelam DK and Kiran M. Fine-needle aspiration cytology of subcutaneous cysticercosis. *Diag Cytopath* 1991; 7(2): 223–224.
- [8]. Liu H, Juan YH, Wang W, et al. Intramuscular cysticercosis: Starry sky appearance. *QJM*. 2014;107(6):459–61.
- [9]. Hawk MW, Shahlaie K, Kim KD, Theis JH. Neurocysticercosis: A review. *Surg Neurol*. 2005;63(2):123–32.
- [10]. Gupta S, Gupta S, Mittal A, et al. A rare manifestation of cysticercosis infestation. *Acta Med Indones*. 2014;46(1):54–57.
- [11]. Tamhane TM, Kachewar SG, Lakhkar DL. Imaging in disseminated cysticercosis: A case report and review of literature. *Indian J Appl Radiol*. 2015;1(1):105.
- [12]. Gupta NK, Panchonia A, Jain D. Cysticercosis of breast. *Trop Parasitol*. 2013;3(2):148–50.
- [13]. Suchitha S, Vani K, Sunila R, Manjunath GV. Fine needle aspiration cytology of cysticercosis – a case report. *Case Rep Infect Dis*. 2012;2012:854704.
- [14]. Sinha S, Tiwari A, Sarin YK, Khurana N. Isolated soft tissue cysticercosis involving the trunk in children: Report of 4 cases. *APSP J Case Rep*. 2013;4(3):35.
- [15]. Venkat B, Aggarwal N, Makhalkar S, Sood R. A comprehensive review of imaging findings in human cysticercosis. *Jpn J Radiol*. 2016;34(4):241–57.
- [16]. Elhence P, Bansal R, Sharma S, Bharat V. Cysticercosis presenting as cervical lymphadenopathy: A rare presentation in two cases with review of literature. *Niger J Clin Pract*. 2012;15:361–63.
- [17]. Chandler AC. *Introduction to Parasitology*. New York, NY: John Wiley & Sons; 1958. pp. 350–54.
- [18]. Jashnani KD, Desai HM, Shetty JB, Pandey I. Fine-needle aspiration cytology of subcutaneous cysticercosis: A series of five cases. *Annals of Tropical Medicine and Public Health*. 2016;9(1):73–75.
- [19]. Chakrabarti S, Bandyopadhyay A, Roychowdhuri A, Mondal S. Incidental diagnosis of cutaneous cysticercosis on cytology: A case report. *International Journal of Medical Science and Public Health*. 2016;5(6).
- [20]. Kraft R. Cysticercosis: An emerging parasitic disease. *Am Fam Physician*. 2007;75:91–98.
- [21]. Lohra S, Barve S, Lohra P, et al. Subcutaneous cysticercosis: Role of high resolution ultrasound in diagnosis. *Natl J Med Res*. 2014;4:82–86.
- [22]. Monteiro L, Almeida-Pinto J, Stocker A, Sampaio-Silva M. Active neurocysticercosis, parenchymal and extra-parenchymal: A study of 38 patients. *J Neurol*. 1993; 241:15–21.
- [23]. Wilson M, Bryan RT, Fried JA. Clinical evaluation of the cysticercosis enzyme, linked immunoelectro transfer blot in patients with neurocysticercosis. *J Infect Dis*. 1991; 164:1007–9.
- [24]. Arora VK, Gupta K, Singh N, Bhatia A. Cytomorphologic panorama of cysticercosis on fine needle aspiration. A review of 298 cases. *Acta Cytol*. 1994; 38:377–80.
- [25]. Suchitha S, Vani K, Sunila R, Manjunath GV. Fine needle aspiration cytology of Cysticercosis - case report. *Case Reports in Infectious Diseases*. Vol 2012, Article ID 854704, 2 pages
- [26]. Kraft R. Cysticercosis: An emerging parasitic disease. *Am Fam Physician*. 2007; 75:91–98.

- [27]. Mittal A, Das D, Aiyer N, et al. Masseter cysticercosis – a rare case diagnosed in ultrasound. *Dentomaxillofac Radiol.* 2008; 37:113–16.
- [28]. Sidhu R, Nada R, Palta A, et al. Maxillofacial cysticercosis – uncommon appearance of a common disease. *J Ultrasound Med.* 2002; 21:199–202.

Lalawmpuii Pachuau, et. al. "Subcutaneous Cysticercosis Mimicking Soft Tissue Neoplasm – A Case Report." *IOSR Journal of Dental and Medical Sciences (IOSR-JDMS)*, 19(8), 2020, pp. 15-20.