

Giant Cell Tumor of Extensor Tendon Sheath in Middle Finger: A Case Report

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Abstract

Giant cell tumor of the tendon sheath is a rare benign soft tissue tumor of hand. Most commonly occur between the third and fifth decades of life at palmar surface. It appears as a painless, perceptible enlargement. It can mimic and make differential diagnoses with several hand tumors. Definitive diagnosis and the treatment of choice are complete resection and histopathological examination. Here we describe a case with clinical presentation like painless, slow growing soft tissue swelling at extensor tendon of middle finger over metacarpophalangeal joint.

Keywords: Giant cell tumor, Extensor tendon sheath, Hand, Excisional biopsy

Journal of Green Life Med. Col. 2021; 6(2): 78- 80

Introduction:

A giant cell tumor of the tendon sheath (GCTTS) is a rare, benign tumor which can develop in the tendon sheaths around the body.¹ It can developed anywhere in the body where there is a tendon sheath, but is most common in the hand and wrist.¹ It is the second commonest tumor of the hand.² Trauma, inflammation, metabolic disease and a neoplastic etiology are considered as etiological factors.³ A GCTTS can occur at any age, but is most common in adults and is more commonly found in women.¹ It is divided into two types based on clinical and biological manifestations: localized and diffuse forms. These can also be intra-articular and extra-articular. The benign localized type affects the hand and fingers, but the more aggressive diffuse form affects major joints.⁴ GCTTS is most commonly found around the index or long finger's distal interphalangeal joint followed by ring and little finger. The following case presents with the features suggesting GCTTS of ring finger.⁵

Case Report:

A 14-year-old boy came to the orthopedic outpatient department, CMH Bogura with a history of painless swelling over metacarpophalangeal (MCP) joint of right middle finger for four-month. The swelling was spontaneous with no history of trauma. It gradually increased in size, hampering the daily living activities of the boy.

On clinical examination, a 1cm x 1 cm firm swelling was seen on the dorsal surface of the right middle finger near MCP joint. During clinical examination there was no localized temperature or tenderness. The swelling was well defined, smooth, firm and uniform in consistency. The swelling was movable sideways with no attachment to the bone and free from over line skin. It adhered to the underlying soft tissue and hence moved with the movement of the finger. Distal neurovascular status was intact. X-ray anteroposterior (AP) and oblique view of right hand showed no abnormality. Ultrasonography was done which revealed mass attached to the underlying soft tissue on the dorsal surface of right middle finger. FNAC from the swelling showed round to oval cells with foci of osteoclastic giant cells with regular nuclei suggestive of a Giant cell tumor of tendon sheath. Other investigations like complete blood counts, chest X-ray, random blood sugar, liver function tests, and renal function tests were normal. The patient was surgically fit. Excisional biopsy of the underlying swelling was performed under local anesthesia. A longitudinal incision was given over the swelling. Soft tissue dissection was done and the tumor

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Received: 01.12.2022

Accepted: 10.12.2022

was seen adhered to the underlying extensor digitorum tendon of middle finger. Excision of tumor was easy and was excised completely. The closure was done in layers. The excised tumor was sent for histopathological examination, which showed polygonal to round histiocytes surrounded by multinucleated giant cells, fibro-fatty tissue suggestive of GCTTS. Postoperative period was uneventful. Hand was mobilized from 2nd postoperative day and stitches were off on 14th post-operative period. The patient was on regular follow up with no functional debility. The patient was on follow up for last six months. The patient was able to perform his activities of daily living with no evidence of recurrence both clinical and radiological examination.

Discussion:

Giant cell tumors of the tendon sheath in the hand are relatively uncommon. Targett first described GCTTS in 1897⁶. The reported age distribution is 8 to 80 years.⁶ Most commonly occur between 30-50 years of age⁷ with a peak incidence in those aged 40-50 years. These tumors are rarely found in patients younger than 10 years or older than 60 years of age. It is more common in female with female to male ratio is 3:2.^{8,9} These tumors may occur anywhere in the hand but more commonly present as firm lobulated masses on the lateral side of index and middle fingers of hand⁶ and commonly seen on the flexor aspect.¹⁰

Tumor classification has an important role to analyzing the recurrence pattern. GCTTS are classified into localized nodular type (common in hand) and diffuse type (common in joints) by Byers.^{11,12} Al Qattan has classified GCTTS into two types, Type-I and Type-II, where Type I- as single tumor which is round or multi lobulated, and Type II, where there are two or more distinct tumors which are not joined together.¹¹

The cause of GCTTS is unknown; however, it might be linked to an inflammatory response, a local lipid metabolic issue, or osteoclastic growth, infection or trauma.¹³ The tumor usually appears as a slowly growing, painless mass on the palmar surface of the fingers.¹⁴

Diagnostic workup includes patient history and a detailed physical examination. Plain radiographs can be helpful since GCTTS may produce erosions in the cortical bone and may invade medullary space. Magnetic resonance imaging (MRI) is the most useful diagnostic tool and is also required for surgical planning. MRI helps to classify GCTTS into type 1 and type 2 according to Al Qattan classification in which type I describes a single round or multilobulated tumor while type II describes two or more distinct, separated tumors.¹⁴

Complete local excision is the treatment of choice for giant cell tumor of the tendon sheath.⁶ The lesions are benign, but recurrence is noted in up to 40% of patients (10% to 20% more commonly reported) even after meticulous excision of the friable fragments.⁶

Risk factors for recurrence or persistence include adjacent degenerative joint disease, location at the finger distal interphalangeal and thumb interphalangeal joint, bony invasion, multifocal disease, tendon involvement, and poor surgical technique⁶. Extensile surgical approaches are frequently required, and gentle blunt dissection should be performed to minimize fragmentation of the encapsulated tumor mass. Magnified vision is helpful to discover discolored synovial tumor, which should be removed during the marginal tumor resection.⁶

Differential diagnosis of hand tumors include lipomas, hemangioma, foreign body, myxoid cyst, synovial carcinoma, tophaceous gout, glomus tumor, tuberous osteitis, epidermal cyst, fibroma, and metastasis.^{15,16}

Although the condition is routinely benign, malignant degeneration has been reported in exceptional cases¹⁶. The diffuse type can be locally aggressive, with reports of possible multiple recurrence and malignant transformation¹⁷. These data reinforce the need for early and accurate diagnosis, and proper treatment.¹⁸

Conclusion:

Our case is an example of GCTTS in a single digit of the hand. The recurrence can be prevented by accurate pre operative diagnosis with FNAC and use of magnification for complete excision with microsurgical skills. Finally, if required, the hand's function should be recreated to minimize the loss.

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