Case Report
A Case of 14-Year-Old Male with Fibroma of Tendon Sheath of the Hand with Novel Chromosomal Translocation 4;10

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Fibroma of tendon sheath (FTS) is an uncommonly encountered soft tissue mass, which is morphologically distinct from the more commonly seen giant cell tumor of tendon sheath (GCTTS). Initially described in 1936, FTS is typically a slow growing, painless, firm mass with a predilection for the upper extremity, frequently involving the hand. Cases of associated triggering or compression neuropathies have been described when underlying tendons or nerves are affected. Currently, the literature on FTS is sparse and largely limited to case reports. More recently, few reports of cytogenetic analysis on FTS have been reported in the literature. Cellular and chromosomal analysis of FTS tissue revealed chromosomal translocations with yet unknown clinical significance.

Here, we present a case report of FTS in a 14-year-old male with a painless enlarging mass of the palmar side of the left hand treated by excision. Subsequent karyotypic analysis revealed a novel chromosomal translocation t(4;10) (p16;q24), add (10)(q22)[24]. To our knowledge, this is the first description of this chromosomal aberration in the literature.

1. Introduction

Fibroma of tendon sheath (FTS) is an uncommonly encountered soft tissue mass, which is morphologically distinct from the more commonly seen giant cell tumor of tendon sheath (GCTTS). Initially described by Geschickter and Copeland in 1936, FTS is typically a slow growing, painless, firm mass with a predilection for the upper extremity [1]. Chung and Enzinger subsequently reviewed 138 cases of such tumors and noted a male to female predilection with a median age of 31 years. The hand and fingers were involved in 81% of the cases with the thumb most commonly affected [2].

The literature on FTS is sparse and largely limited to case reports or small series [3–7]. Commonly, patients present with a painless, minimally tender solitary mass. However, cases of associated triggering or compression neuropathies have been described when underlying tendons or nerves are affected [2–7]. Recurrence rates vary and, although rare in many case reports, have been recorded up to 24% in larger series [2–6].

More recently, few reports of cytogenetic analysis on FTS have been reported in the literature. One case of FTS identified a clonal chromosomal abnormality t(9;11) (p24;q13-14), while another identified a cellular abnormality t(2;11)(q31-32;q12) in half the cells sampled [8, 9]. The clinical significance of these translocations has yet to be fully elucidated.

Here, we present a case report of FTS in a 14-year-old male with a painless enlarging mass of the palmar side of the left hand treated by excision. Subsequent karyotypic analysis revealed a novel chromosomal translocation t(4;10) (p16;q24), add (10)(q22)[24].

2. Case Report

A 14-year-old right-hand-dominant Caucasian male presented with a history for several years of a slowly enlarging mass of the left palm centered about the third metacarpal head (Figure 1). The mass was not associated with pain except when batting during baseball. He denied any sensory
deficits or motor weakness. Physical examination demonstrated a soft, fixed mass about the palmar aspect of the left hand extending from the region of the second to the fourth metacarpal heads. His neurologic assessment was unremarkable.

Plain radiographs did not show any abnormalities. Magnetic resonance imaging (MRI) with intravenous gadolinium demonstrated a 3.6 × 1.6 × 2.4 cm well-defined soft tissue mass centered about the third metacarpophalangeal joint abutting the flexor tendon sheath of the third digit (Figure 2). The mass demonstrated mild enhancement with contrast. There was no osseous erosion or joint effusion noted. Based on this, the differential diagnosis included infectious etiology versus soft tissue mass including giant cell tumor of tendon sheath, lipoma, leiomyoma, and neuroma.

The patient underwent excisional biopsy under general anesthesia with tourniquet control after gravity exsanguination. An incision was made in the distal palmar crease overlying the mass, and soft tissue dissection was carried down through the level of the palmar fascia. Intraoperatively, a solitary, solid tan-colored mass was encountered. It was well-encapsulated without communication to any neurovascular or tendinous structure aside from the A1 pulley of the third flexor sheath (Figures 3 and 4). The lesion was removed en bloc. An intraoperative frozen section was performed. The tissue was considered lesional and was deferred for examination by an orthopaedic pathologist.

Final pathologic examination revealed a 3.8 × 2.5 × 1.3 cm spindle-cell lesion with mild lobulation. The spindle cells resembled fibroblasts, and no nuclear atypia was seen. The lesion was relatively hypocellular without necrosis or
hemorrhage. Multinucleate giant cells were not seen (Figure 5). The tumor cells demonstrated focal reactivity to CD34 and smooth muscle actin. Rare cells demonstrated immunoreactivity to calponin. The tumor cells were negative for CD31, Fli-1, CD68, muscle-specific actin (HHF-35), and desmin. Less than 1% of the cells demonstrated proliferation via Ki-67, consistent with a benign process. The histologic and immunohistochemical findings were mostly in keeping with fibroma of tendon sheath. In addition, a portion of the fresh specimen was submitted for karyotype analysis. The results revealed a 46,XY,t(4;10)(p16;q24), add (10)(q22) in twenty-four analyzed cells. All cells analyzed showed a translocation involving chromosomes 4 and 10 and a complex structural rearrangement of the long arm of the other chromosome 10 at band q22.

The patient’s postoperative course was uneventful, and he returned to baseball activities 6 weeks postsurgery. He is currently at 21-month status postsurgical excision, reports pain-free activity, and exhibits no evidence of local recurrence (Figure 6).

3. Discussion

FTS is a relatively uncommon tumor that typically presents as a painless, slow growing, firm nodule in the palm or digits. Typically, FTS is seen in the young to middle-aged population, with some series reporting an up to 3:1 male to female ratio [10]. Uncommonly, antecedent trauma is reported [10]. The differential diagnosis should include giant cell tumor, mucinous cyst, lipoma, synovial sarcoma, epidermal cyst, and leiomyoma [3]. Clinically, FTS most commonly behave like giant cell tumors of tendon sheath in that they are firm masses with an intimate anatomic relationship to the tendon sheath. They additionally exhibit similar signal patterns on MR imaging with low to intermediate signal intensity on

![Figure 5](image1.png)

**Figure 5:** (a) Low-power examination demonstrates a mildly lobulated lesion with tumor cells set in a collagenous matrix. The lesion shows an admixture of hypocellular and hypercellular areas (hematoxylin and eosin stain, magnification 10x). (b) High-power examination demonstrates fibroblasts with basophilic nuclei in a focally hypercellular area. No mitotic figures are identified (hematoxylin and eosin stain, magnification 40x).

![Figure 6](image2.png)

**Figure 6:** (a, b) Clinical postoperative images of patient 21 months after surgery.
References
