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Citation: Kar Sk, Recurrent Fibrosarcoma Protuberans of Anterior Chest and Abdominal Wall. A Rare Case Report. Arch Med. 2015, 8:1.

Abstract
Dermatofibrosarcoma Protuberans (DFSP) is a locally invasive neoplasm of soft tissue. Though distant metastasis is rare, it has high recurrence rate. The authors here report a case of recurrent DFSP in a 50 year male who presented with a large protuberant tumor involving anterior chest and abdominal wall with a history of fleeting pain. Wide local excision was performed with 3 cm free margin followed by repair of the wound with a right latissimus dorsi flap and a non-absorbable mesh. Histopathology and immunohistochemical staining findings were consistent with the diagnosis of dermatofibrosarcoma protuberance.

Introduction
Dermatofibrosarcoma Protuberans (DFSP) is a rare soft tissue tumour. It often has a high recurrence rate. Commonly the tumor involves the trunk (42-72%), proximal extremities (16-30%) and the head and neck (10-16%) [1-6]. However, it can metastasize in 1-4% of cases after so many years of development of primary lesion [7]. It is a low to intermediate grade malignancy in 90% cases [7] and rarely progresses to high grade fibrosarcomatous component [8]. The incidence rate is only 0.8 cases per million with slight male predominance [9]. Usually middle aged males suffer more from this tumor as compare to female [10]. Recently the authors have encountered such a case of recurrent DFSP which was treated successfully.

Case Report
A 50 year male presented with a large swelling involving the anterior chest wall and the anterior abdominal wall with occasional pain and local irritation. Patient has a burn injury at his age of 4 years and developed scar tissue over anterior abdominal and chest wall. After few years he noticed a small firm painless swelling over the scar which progressively increased in size and patient developed symptoms of pain for which FNAC followed by excision of mass was done. Histopathology and immunohistochemical staining findings were consistent with the diagnosis of DFSP.

Recurrent Fibrosarcoma Protuberans of Anterior Chest and Abdominal Wall. A Rare Case Report

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Discussion

Less than 5% of all thoracic tumors is malignant [11] of which 50% is soft tissue fibrosarcoma. DFSP accounts for only 2-6% of all soft tissue sarcomas. Darier and Ferrand first described DFSP as a distinct cutaneous disease and termed as progressive and recurrent dermatofibroma in 1924 [12]. However it was initially published in literature as early as 1890. In 1925 Hoffman coined it as “dermato-fibrosarcoma-protuberans”. The origin of DFSP is still controversial (Figure 6). With this defect usually genetic correlation is associated. This occurs due to translocation of chromosomes 17 and 22 or formation of a ring [13]. Primarily the tumor starts with a very small non indurated nodule and patient presents when it slowly increases to large, firm, sometimes tender, multilobulated, blue to red mass as in our case. Due to its local aggressiveness, it can grow into the surrounding fat, fascia, muscle and even bone. In this case, tumor invaded the lower part of sternum and adjacent four ribs bilaterally. So, during mass removal, these bones were excised in this case. Rarely, distant wall was reconstructed. Biopsy revealed Dermatofibrosarcoma Protuberans with equivocal margin. Immunohistochemical staining was diffusely positive for CD34 and negative for CD117. Post operative course was uneventful. Postoperative chemotherapy for mitotic activity and radiotherapy for equivocal margin were advised by the oncologist. Patient was discharged on 10th post operative day and advised monthly follow up.
metastasis is seen in such case and it is nearly 4-6%, with the lungs being the common site [14,15]. There was no distant metastasis in this patient. This case was a recurrence of primary tumor.

Literature shows that DFSP has characteristically high and varying recurrence rate ranging 10-80% [16]. Surgery is the gold standard treatment. Excision of mass with 3 cm tumor free margin is the mainstay though histological free margin as local control is emphasized [17]. In this case along with muscle flap, non absorbable mesh was used to reinforce the anterior abdominal and chest wall to avoid post-operative complication of incisional hernia.

Chemotherapy is not always advised but radiotherapy is usually recommended if tumor margin is equivocal. In this case, chemotherapy was advised for mitotic activity of this aggressive tumor and also post operative radiotherapy was recommended as adjuvant therapy to reduce the chance of recurrence when excision biopsy reveals doubtful tumor margins [18].

In the era of minimally invasive surgical practice, a microscopic surgical procedure termed as Mohs surgery is widely popular not only for its high oncological effectiveness, it can also provide maximal tissue salvage. In case of unresectable tumor tyrosin kinase inhibitor (Imatinib) is the treatment of choice. Among all available treatment of such case, primary surgical excision is the gold standard and early resection with tumor free margin plays the pivot role of success in the management of DFSP. It provides advantage of less chance of recurrence and sometime needs no chemotherapy or radiotherapy, thus enhances the positive outcome with good prognosis.

Conclusion
DFSP is a locally invasive tumor with rare incidence of distant metastasis though recurrence rate is high. Early detection and wide primary surgical resection followed by postoperative adjuvant radiotherapy and chemotherapy can successfully prevent recurrence resulting a favorable outcome.
References


