Primary tumours arising in sternum are extremely rare. We came across two patients having primary tumours arising from sternum, evaluated them for clinical diagnosis and treated them surgically by performing wide local excision and reconstruction. Histopathological examination of the excised specimens revealed one of them to be Ewing’s sarcoma and the other giant cell tumour. These tumors are very uncommon as evidenced from published English literature. We adopted two different techniques of reconstruction in both the cases based on the need and feasibility. We consider both the reconstructive techniques to be novel and ideal for such situation. After a six month follow up both the patients were doing well in terms of function and aesthetics of the operated region.

Case Reports

Case 1

A 16 years old boy presented with a swelling over the sternal region noticed over a period of six months. The lesion was subjected to incisional biopsy 1 month before the patient attended the outpatient department of our hospital. The patient was evaluated by our team of thoracic surgeons and the plastic surgeons and treatment planned. The previous incisional biopsy showed none of the features of malignancy. Hence the treatment was planned considering it a benign tumour. MRI depicted the lesion expanding into the mediastinum and externally as well. The tumour was found to involve medial ends of both the clavicles and sternal end of the upper two ribs.

Wide local excision was performed including the tumour with 2 centimeters of healthy margin from sternum, clavicles and the ribs. The tumour was free from adhesion with the mediastinal tissues and could be taken out with ease and it was removed with its overlying skin. Hence the excision left a defect of significant size with the mediastinum and the pericardium exposed to the exterior without any tissue overlying them (Figure 1).

From the reconstructive point of view the bony wall of the thorax along with overlying soft tissue and skin were deficient. The part was planned to be reconstructed with rib grafts with a cover of pectoralis major muscles and a split thickness skin graft over it. An inframammary approach was used to harvest the

**Figure 1:** Ewing’s Sarcoma (1-Rib graft, 2- Pectoralis Major muscle cover, 3 - After complete healing, 4 - 400X microscopic view after H & E staining).
muscle flap as well as the ribs. The ribs were fixed with titanium miniplates (2 mm) and cross head titanium screws (2x10 mm). Reconstructed bony wall was covered with pectoralis major muscle and split thickness skin graft was put over that. The recovery was uneventful. Post excisional histopathology study established the diagnosis of Ewing’s sarcoma. The patient was referred to the department of medical oncology for adjuvant chemotherapy. He was observed doing well after 6 months of surgery and completion of chemotherapy.

Case 2

A 32 years old female presented with a swelling over sternal region observed by the patient over a period of 8 months. It was growing slowly. She was operated for giant cell tumour of lower end of left radius 2 years prior to this observation. Radiographic evaluation suggested it to be a giant cell tumour of sternum. The tumour was expanding into the mediastinum with very minimal growth towards the surface. The patient was followed up for six months and not found to develop any complication. She probably had the necrosis or atrophy of medial part of pectoralis muscle covering it.

Considering the size of the bony defect we preferred to proceed for allo-implant. Titanium mesh was used for reconstruction of bony defect, covered with pectoralis major muscle flaps from both the sides. Skin wound could be closed primarily without any difficulty. The patient was followed up for six months and not found to develop any complication. She probably had the necrosis or atrophy of medial part of pectoralis muscle flap which was evident from the contour of the reconstructed area during follow up (Figure 2).

Discussion

Primary tumours of sternum are very rare [1]. The bone is mostly affected secondarily due to metastasis from other malignant tumours like breast, ovary, kidney etc [2]. Ewings sarcoma is the second most common primary malignancy of sternum [3]. Very few cases of primary Ewings sarcoma and primary giant cell tumour arising in sternum have been reported. The sternal reconstruction is necessary in order to achieve protection to the exposed thoracic viscera and aesthetic appearance of the chest wall. Available materials and techniques are plenty to choose [4]. It includes metallic/alloplastic prostheses, autogenous bone graft, cryo–preserved sternal allograft or homograft as skeletal substitutes which need to be covered with soft tissue like pectoralis major or latissimus dorsi, rectus abdominis muscle or myocutaneous flap or omental flap [5,6]. Rib grafts and pectoralis major muscles we used in first case for reconstruction, are in close proximity of these defects and can easily be obtained by any reconstructive surgeon. Bone banks are available only in select healthcare delivery centers. Titanium mesh is well tolerated without much of complications and suitable for large sternal defects [7]. Titanium mesh we used in the second case was essential due to size of the defect. The techniques we used for reconstruction of post excisional defects are technically reproducible, cost effective and bio-compatible. Both the options used for reconstruction do not pose any threat in terms of adjuvant radiation or chemo therapy. The negativity of incisional biopsy in the case of Ewings sarcoma of course was an issue considering retrospectively since neo-adjuvant chemotherapy is recommended for such cases before contemplating to surgical excision. We presume it to be a procedural inadequacy in part of the surgeon performed this.

Disclosure

We disclose no conflict of interest in these cases and no financial support is received from any source for these cases.

References

1. Eng J, Sabanathan S, Pradhan GN (1989) Primary sternal tumours. Scand J Thorac Cardiovasc Surg 23: 289-292. Link: https://tinyurl.com/y6f5q7f

2. Bandi V, Lunn W, Ernst A, Eberhardt R, Hoffmann H, et al. (2008) Ultra-sound vs. CT in detecting chest wall invasion by tumor: a prospective study. Chest 133: 881-886. Link: https://tinyurl.com/y4dvyy9d

3. Delattre O, Zucman J, Melot T (1994) The Ewing family of tumors – a subgroup of small-round-cell tumors defined by specific chimeric tran-scripts. N Engl J Med331: 294-299. Link: https://tinyurl.com/y4la9san

4. Sanna S, Brandolini J, Pardolesi A, Argnani D, Mengozzi M, et al. (2017) Materials and techniques in chest wall reconstruction: a review. J Vis Surg 3: 95. Link: https://tinyurl.com/y4k7yz8f

5. Nosotti (2012) Sternal reconstruction for unusual chondrosarcoma: Innovative technique. J Cardiothorac Surg 7: 40. Link: https://tinyurl.com/y3ar8mzh

6. Seder CW, Rocco G (2016) Chest wall reconstruction after extended resection. J Thorac Dis 8: S863-S871. Link: https://tinyurl.com/y6fupfpf

7. Zhang Y, Li JZ, Hao YJ (2015) Sternal tumor resection and reconstruction with titanium mesh: a preliminary study. Orthop Surg 7: 155-160. Link: https://tinyurl.com/y48fnpj5

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