Myxofibrosarcoma of Base of Skull in Paediatric Age Group: A Case Report

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Abstract
Myxofibrosarcoma is a common sarcoma affecting extremities of elderly age group. It rarely affects the younger age group and the head neck region. It exhibits an infiltrative growth pattern and has a high local recurrence rate. We report a 14 year old patient with a myxofibrosarcoma of base of skull. The patient’s age and clinical presentation favored a bedside diagnosis of a dermoid cyst or langerhans cell histiocytosis, radiology was suggestive of an intraosseous hemangioma and surgical pathology favored a low grade osteosarcoma. However, tissue immunohistochemistry clinched the diagnosis of a Myxofibrosarcoma. He underwent a near total excision with adjuvant radiotherapy and remains recurrence free at one and a half year follow up. Tissue immunochemistry played a pivotal role in the management of our patient. We have attempted to bring forward the pertinent neurosurgical issues related to the management of this common soft tissue sarcoma which is rarely encountered by Neurosurgeons.

Introduction
Soft tissue sarcomas are a heterogeneous and a diverse group of neoplasm of mesodermal origin. Myxofibrosarcoma is one of the common histologic types of Soft Tissue Sarcomas. Formerly myxofibrosarcoma was considered as the myxoid variant of Malignant Fibrous Histiocytoma. The 2002 WHO classification of tumors included them as a distinct entity after Angervval & kindblom [1] recognized them to be so and coined the term Myxofibrosarcoma. To date less than 20 patients have been reported to have head neck Myxofibrosarcoma and ours is the first case of a skull base myxofibrosarcoma in the pediatric age group.

Case History

Figure 1: Preoperative lateral and posterior profile view depicting the left occipital swelling.
A 14-year-old boy presented with a one-year history of a painless, progressive swelling over the left occipital region (Figure 1). He denied constitutional symptoms and was neurologically intact on examination. Contrast enhanced computed tomogram (Figure 2) head revealed a large, hyper vascular, expansile lytic mass of the left occipital bone. It was well corticated with occasional bony internal septations. It was heterogeneously hyper intense and showed intense post contrast enhancement. The basal cisterns were effaced with significant mass effect on brainstem and cerebellum. An angiogram revealed retention of contrast on delayed phase images suggesting a possibility of intraosseous hemangioma of occipital bone. Intraoperatively, the lesion was pearly white and fleshy in consistency. Outer and inner bone tables were paper thin. The bone tables were breached but the dura and scalp was uninvolved. He underwent a near total excision, with the nubbins infiltrating jugular foramen being left behind. Postoperatively, the patient was clinically stable, with a transient lower cranial nerve paresis which recovered gradually (Figure 3).

Pathological evaluation of the surgical specimen revealed a low grade Osteosarcoma. Immunohistochemistry clinched the diagnosis of a myxofibrosarcoma (Figure 4). He received adjuvant radiotherapy in view of the residual disease (Figure 5) and the IHC findings. At 30 months follow up he remains recurrence free.
Discussion

Myxofibrosarcoma is the most common sarcoma in the elderly age group. Its peak incidence is in the seventh decade with Slight male Preponderance. It involves the extremities in over 75% cases, trunk, retroperitoneum and mediastinum being the less favored sites. It exhibits a slow and painless progression [2]. Based on the plane of origin Mentzel et al classified myxofibrosarcoma into two subgroups: superficial (dermal/subcutaneous) and deep (intramuscular/sub fascial) [3]. The superficial group of tumors exhibit a longitudinal pattern of extension and infiltration, whereas the intramuscular/sub fascial tumors form a single discrete mass.

These tumors have been subdivided into three or four grades based on the degree of cellularity, nuclear pleomorphism and mitotic activity [3,4]. Mutter et al. [5] have reported a local recurrence rate of 14.3% for Myxofibrosarcoma [5]. Other groups have reported a high Local recurrence rate of upto 61% [1,3,6]. An infiltrative growth tendency with ability to extend along facial and vascular plane may be implicated in the high local recurrence rates found in these tumors [3-5,7].

The tendency of myxofibrosarcoma to be of progressively higher grade in recurrences by becoming gradually more cellular, more pleomorphic, more mitotically active, and more necrotic seems to be a distinctive feature and is in contrast to most other sarcomas [3].

Imaging

On Magnetic resonance imaging Myxofibrosarcoma exhibit abnormal signal infiltration along fascial planes. Presence of a tail sign is common and associated with an adverse local recurrence free survival [8]. Ahmadi et al. [9] have proposed that a fairly uniform dural enhancement in vicinity of a neoplasm on Contrast MRI is most likely reactive but focal breach in continuity of enhancing adjacent dura implies infiltration and warrants dural excision, where possible [9,10].

Treatment

A wide margin surgical resection aimed at achieving negative margins is the main modality of treatment of Myxofibrosarcomas. Radiation is an effective adjuvant treatment in these patients and should be considered in the local management of this disease [5]. The need for follow-up imaging to assess tumor recurrence or progression of residual disease makes selection of materials used in closure of the cranial defect after cranial surgery a key element of operative planning. Poly ethyl ethyl ketone and titanium implants produce minimal metallic artifact on MRI and CT, making it an excellent option for cranial reconstruction when autologous bone is unavailable or not suitable.

Treatment Data to guide the optimal management strategy for Myxofibrosarcoma of the calvaria and skull base is sparse.
Myxofibrosarcoma of skull present a challenging management conundrum owing to the combination of exceedingly rare occurrence, varied pathologic features, infiltrative tendency, high local recurrence rate and stage progression in recurrences. Currently ensuring a tumor free margin, an expert pathological review, adjuvant radiotherapy and a close surveillance form the cornerstone of treatment plan. The future of management lies in understanding the molecular basis of the tumors to develop and direct therapy at them.

References