Mandibular myofibrosarcoma of childhood: Surgical resection & reconstruction with fibula flap

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INTRODUCTION

Myofibrosarcomas are rare soft-tissue sarcomas mainly affecting the head and neck or extremities of adult patients. The evolution of the cell of origin, that is the myofibroblast, still eludes us. Described as a modified common fibroblast, it is either a native constituent of normal tissues, or the result of tissue reactive processes.

The natural course of myofibroblastic malignancies follows a slowly-growing, infiltrative pattern of spread, but carries a risk of recurrence and metastasis, even after many years. The latter attribute can be explained by the lack of a capsule.

Below, we report a case of a mandibular myofibrosarcoma in an 11-year-old girl.

CASE PRESENTATION

An 11-year-old girl presented with a left-sided mandibular tumor. Initially, it had been enucleated transorally, before the patient was evaluated in our clinic. Histopathology was consistent with intermediate-to-high grade myofibrosarcoma. A CT scan one month post-operatively demonstrated a residual mass, measuring 3.7cm transverse x 1.9cm antero-posterior x 3.4cm cephalo-caudal, and involving the majority of the ramus.

Segmental mandibulectomy was performed, encompassing the angle of the jaw (Fig. 1A). An osseofasciocutaneous fibula flap was harvested, including 10cm of bone. The straight fibular segment, prior to its incorporation into the defect was properly osteotomized to resemble the shape of the jaw angle (Fig. 1B). The soft-tissue portion of the flap was inset intraorally to cover the fibular bone. In addition, it would be used as a monitoring paddle. Before closure of the donor site, orthopedic surgeons performed osteosynthesis of the distal fibula to the tibia, to secure the stability of the ankle. The pathology report was consistent with myofibrosarcoma (Fig. 1C). Six months after the procedure, there is no evidence of recurrence or metastasis. In addition, the patient demonstrates acceptable occlusion and speech quality. No donor site complications were noted, and the child gradually resumes the physical activities of her peer group.

DISCUSSION

Malignancies showing myofibroblastic differentiation are very rare and their biologic behavior is far from being elucidated. Myofibrosarcomas are soft-tissue sarcomas but a few of them are intraosseous, as the case with our patient. Myofibroblasts (Fig. 1C), which give rise to these sarcomas, are reactive cell types, induced by trauma, fibrosis and adjacent tumors. None of the above situations was noted in our patient. Interestingly, the normal periodontal ligament is amputated in the very few sites in which a native myofibroblastic population has been detected, but the nidus of our patient’s neoplasm was posterior to the tooth-bearing mandible.

Myofibrosarcoma is a rare and undefined entity, so that every new report of a case might contribute to understanding its natural course and management. In view of all the previous reports, the treatment of choice should be surgical resection. Here, we presented a case in a pediatric patient, trying to emphasize the importance of a comprehensive approach to a superior oncological outcome, as well as acceptable quality of life.

REFERENCES

1. Mertens T, Dry S, Katenkamp D, Fletcher CD. Low-grade myofibroblastic sarcoma: analysis of 18 cases in the spectrum of myofibroblastic tumors. Am J Surg Pathol. 1998;22(10):1228-38.
2. Fisher C. Myofibrosarcoma. Virchows Arch. 2004;445(5):215-25.
3. Smith DM, Mahmoud HH, Jenkins JJ, 3rd, Rao B, Hopkins KP, Parham DM. Myofibrosarcoma of the head and neck in children. Pediatr Pathol Lab Med. 1995;15(3):403-18.
4. Nattan SS, Hung-YI, Dia I, Altharaisen E, Boland P, Cordeiro PG, et al. Ankle instability after vascularized fibular harvest for tumor reconstruction. Ann Surg Oncol. 2007;14(3):57-64.
5. Baughan B, Demijnin A, Leveque GY, Lapalme-Chaput L. The pattern of facial growth before and during puberty, as shown by French-Canadian girls. Ann Hum Biol. 1979;6(1):59-76.
6. Upton J, Guo L, Labow BI. Pediatric free-tissue transfer. Plast Reconstr Surg. 2009;124(6 Suppl):e313-26.

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