CASE REPORT

Maxillary mesenchymal chondrosarcoma leading to a diagnosis of Li-Fraumeni syndrome

Eduardo Ventura1,* , Sílvia Dionísio1, Ângela Ferreira1, Rute Saleiro1, Hugo Marques1, Manuel Magalhães2, and Carlos Monteiro1

1Maxillofacial Surgery Unit, Centro Hospitalar e Universitário do Porto, Porto, Portugal, and
2Oncology Department, Centro Hospitalar e Universitário do Porto, Porto, Portugal

*Correspondence address. Maxillofacial Surgery Unit, Centro Hospitalar e Universitário do Porto, Largo do Prof. Abel Salazar, 4099-001, Porto, Portugal.
Tel: (+351) 222 077 500; Fax: (+351) 22 2077595. E-mail: simoesventura@gmail.com

Abstract

Mesenchymal chondrosarcoma (MCS) is a rare histological variant of chondrosarcoma, with aggressive behaviour. Due to the unique nature of this disease, management strategies are not well established. Li-Fraumeni syndrome (LFS) is a rare cancer predisposition syndrome with a wide tumour spectrum, associated with TP53 germline mutations. We report a case of MCS of the maxilla, treated with surgical excision and adjuvant chemotherapy, in a patient with a past medical history of choroid plexus papilloma and a family history of early age first-degree cervical uterine cancer, that led to the clinical suspicion of a cancer predisposition syndrome and the subsequent diagnosis of LFS. This is the first MCS described in a LFS case. It demonstrates that adjuvant chemotherapy should be considered, in conjunction with surgical excision, in MCS and that cancer predisposition syndromes should be suspected in patients with multiple neoplasms and a strong family history of cancer.

INTRODUCTION

Mesenchymal chondrosarcoma (MCS) is a rare histological variant of chondrosarcoma that is mostly seen in young adults, unlike conventional chondrosarcomas. It usually presents aggressive behaviour with an inclination for late recurrence and metastasis. Due to the rare nature of this disease, management strategies are not well established, but extensive surgical resection is unanimously described as the mainstay of treatment [1]. Nevertheless, due to the scarcity of clinical cases and data in the literature, the effectiveness of adjuvant chemotherapy and radiotherapy in addition to an ablation strategy is not well defined.

Li-Fraumeni syndrome (LFS) is an aggressive cancer predisposition syndrome, with a broad tumour spectrum, associated with TP53 germline mutations. The neoplasms reported most often are soft tissue sarcomas, osteosarcoma, premenopausal breast cancer, brain tumours and adrenocortical carcinomas [2].

CASE REPORT

A 35-year-old male patient was referred to our department, complaining of a painless, rapid growing exophytic gingival mass in the maxilla with two months’ duration. He had a previous history of choroid plexus papilloma, treated with surgery and radiosurgery at the age of 23. His mother was diagnosed with cervical uterine cancer at the age of 38.

On examination, a lobulated, firm mass was seen extruding from the anterior maxillary vestibule (Fig. 1). All maxillary teeth
were painless to percussion, and normal tooth mobility was present. No cervical lymphadenopathy was detected, nor any changes with cranial nerve examination. A maxillofacial computerized tomographic (CT) scan revealed an osteolytic lesion on the maxillary alveolar process (Fig. 2).

We performed an incisional biopsy of the mass that revealed findings suggestive of a preliminary diagnosis of MCS. After discussion in a multidisciplinary Head and Neck Oncology meeting, the team decided that surgery and adjuvant chemotherapy was the most appropriate treatment plan.

Surgical access was obtained intraorally, and an en bloc resection of the premaxilla was performed (Fig. 3). Histopathological examination confirmed the diagnosis, revealing a completely resected MCS with clear margins (Fig. 4).

The patient underwent an adjuvant chemotherapy regimen with vincristine, doxorubicin, cyclophosphamide/ifosfamide and etoposide (VAC/IE), completing 12 cycles uneventfully. He was referred to the Medical Genetics Department, where further studies confirmed a pathologic variant on the TP53 gene and the diagnosis of LFS.

With 34 months of follow-up, the patient is asymptomatic without any local signs of recurrence, and a good functional and aesthetic result was accomplished with a removable partial denture (Fig. 5). At present, he is in a strict cancer surveillance regimen based on the NCCN protocol.

**DISCUSSION**

MCS is a rare histological variant of chondrosarcoma, mostly seen in young adults, unlike conventional chondrosarcoma that shows a propensity for older age groups. When present in the head and neck area, this condition shows a predisposition for the maxillary anterior alveolus, due to the presence of nasal cartilage [1].

A study of 36 MCS of the maxilla demonstrated that the most common clinical presentation was a mass or growth in 68% of cases, followed by nasal obstruction (32%), epistaxis (32%) and tooth mobility (24%) [3]. In our report, the patient’s chief complaint was a rapid protruding mass that possibly due to its exophytic nature was not associated with nasal symptoms or tooth mobility.

Diagnosis of MCS can only be established by histopathological examination, characterized by a biphasic pattern, with a sheet-like or patternless proliferation of small differentiated spindle or round cells surrounding discrete nodules of differentiated hyaline cartilage [3].
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Figure 4: The typical biphasic pattern of MCS, composed of islands of cartilage (∗) admixed with poorly differentiated small round cells with scant cytoplasm (#). H&E stain.

Figure 5: Rehabilitation with a removable partial denture (A). Frontal view after oral rehabilitation (B).

Extensive surgical resection is described unanimously as the mainstay of treatment but, due to the scarcity of clinical cases and data in the literature, the effectiveness of adjuvant chemotherapy and radiotherapy is not well defined, and consensus has not been reached on the optimal management of MCS [1]. In this case report, the patient received a VAC/IE Adjuvant Chemotherapy scheme, in compliance with the NCCN Clinical Practice Guidelines [4] with excellent tolerability and minimal side effects.

In our case, the rare nature of MCS, the history of choroid plexus papilloma and the history of the patient’s mother developing uterine cancer at the age of 38 lead us to the suspicion of a cancer predisposition syndrome that was confirmed by our Medical Genetics Department.

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CONFLICT OF INTEREST STATEMENT

We have no conflicts of interest.

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