CASE STUDY

Extramedullary Plasmacytoma of the Larynx. A Case Report

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Abstract Extramedullary plasmacytoma of the larynx and localised laryngeal amyloidosis are two entities that are extremely rare in children. We report the case of an 11-year-old child presenting with progressive dysphonia, with a diagnosis of extramedullary plasmacytoma and localised laryngeal amyloidosis. The treatment he received and subsequent follow-up were compared with the few cases found in the literature.

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PALABRAS CLAVE
Plasmocitoma extramedular; Amiloidosis; Plasmocitoma laríngeo; Población pediátrica

Plasmocitoma extramedular de laringe. A propósito de un caso

Resumen El plasmocitoma extramedular de laringe y la amiloidosis localizada laringea son dos entidades infrecuentes y extremadamente raras en niños. Presentamos el caso de un niño de 11 años con disfonía progresiva, diagnosticado de plasmocitoma extramedular y amiloidosis localizada laringea. Comparamos su tratamiento y posterior evolución con los escasos casos encontrados en la literatura.

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Clinical Case

We present the case of an 11-year-old boy sent to the reference hospital due to dysphonia of 2 months’ evolution, without respiratory and deglutitive disruptions.

Physical examination showed a right hemilaryngeal tumour that bulged towards the glottal lumen, reducing ipsilateral mobility. We were unable to define its extent to the right vocal fold (RVF) and the remainder of the ENT exploration was normal (Fig. 1A). The MRI revealed a submucosal glottic tumour centred on the RVF, which deformed the right aryepiglottic fold and obliterated the ipsilateral pyriform sinus, discarding vascular aetiology (Fig. 2A).

We performed laryngeal microsurgery, with intraoperative amyloidosis biopsy and reduction of the tumour mass with CO\textsubscript{2} laser.
Extramedullary plasmacytoma is an uncommon entity, which is extremely rare in children as its mean age of presentation is 56–59 years,1 with a predominance in males. We have found only one case of coexistence of amyloidosis and laryngeal plasmacytoma in children described in the literature: a 12-year-old girl, published by Nagasaka et al.2

The symptoms are caused by local growth, with hoarseness, dysphagia, cough and dyspnoea being the main manifestations. Pribitkin et al.,3 in a study in patients with laryngeal amyloidosis, pointed to hoarseness as the main manifestation, while Riazza et al.4 pointed to dyspnoea.

The diagnosis of extramedullary plasmacytoma is fundamentally histological, based on the presence of plasma cells showing apple-green birefringence with Congo red staining, presenting cytologically benign traits; however, immunohistochemical study shows monoclonality, pointing to their neoplastic nature.5 In our case, immunohistochemical techniques expressed lambda chains, with a negative expression of kappa chains, and the molecular study showed B lymphoid monoclonality.

An extension study is required to establish a differential diagnosis with multiple myeloma and solitary bone plasmacytoma, considering that extramedullary plasmacytoma is generally a localised entity associated with long survival.5

In the treatment of laryngeal amyloidosis, excision of the lesion with CO₂ laser is effective because, despite having invasive capacity, altering the functionality of the vocal fold, the lesion behaves like a benign tumour and does not usually present recurrence in localised forms.1 Nevertheless, some authors advocate an expectant attitude, reserving surgery for specific cases.6

Extramedullary plasmacytoma is highly radiosensitive,7 with local control being achieved in 95% of cases with total doses of 40–60 Gy.8 In the patient described, we excised the lesion with CO₂ laser in the first surgical step, obtaining significant clinical improvement and reduction of the tumour mass. After the definitive AP report of extramedullary plasmacytoma, we decided to apply radiotherapy, achieving the eradication of the plasmacytoma, although the amyloid mass persisted. Given the patient’s limited symptoms, we chose to conduct periodic reviews, observing good clinical evolution.
Conclusion

- The presence of localised laryngeal amyloidosis and laryngeal plasmacytoma could be the two infrequent entities in children.
- The diagnoses are mainly histological and immunohistochemical.
- Treatment should be aimed at eradicating the plasmacytoma and reducing the mass effect. However, in some cases with few symptoms, it may be limited to regular controls.

References


