Case Report

Hemangiopericytoma of the Parapharyngeal Space: A Diagnostic Challenge

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Key Words
Parapharyngeal space · Hemangiopericytoma · Vimentin · MRI · CT scan

Abstract
Parapharyngeal space tumors are known for having a difficult approach, misleading diagnosis and for representing a treatment challenge. Hemangiopericytomas account for less than 1% of all vascular neoplasms and 3% of all soft tissue sarcomas. Only 14 cases have been reported in the worldwide literature in this location. We present a case of a 44-year-old male who was referred for evaluation. A CT scan and MRI showed a large parapharyngeal mass of a possible salivary gland origin. The patient underwent a lateral cervicotomy associated with a transparotid-transmandibular approach, obtaining a vimentin-positive immunostaining tumor defining the diagnosis. The accurate management and prognosis of this type of neoplasm are provided by the definite diagnosis obtained by a correct histopathologic assessment. A high clinical suspicion is essential.

Introduction

The parapharyngeal space located deep within the cervical region is divided in the prestyloid and retrostyloid compartments. Most of the salivary gland neoplasms are located in the prestyloid space, and neurogenic tumors account for most of the retrostyloid masses [1]. Tumors in this area are known for having a difficult approach, misleading diagnosis and for representing a treatment challenge. The clinical course is usually asymptomatic even when tumors reach large sizes. Some symptoms may include dysphagia, otalgia, dyspnea and
foreign body sensation [2]. Imaging studies are essential for evaluation due to the limitations of physical examination.

Hemangiopericytoma (HPC) is a vascular tumor that accounts for less than 1% of all vascular neoplasms and 3% of all soft tissue sarcomas [3]. In the head and neck area, the most common sites include the jaw, parotid gland, nasal cavity, orbit, masticator space, paranasal sinuses and the jugular foramen [3].

Case Report

A 44-year-old male with no significant medical history was referred for the evaluation of a parapharyngeal left cranial base asymptomatic tumor that was found incidentally during a sleep apnea study. A cervical MRI revealed a parapharyngeal mass of 5.7 × 4.6 × 2.8 cm of a possible salivary gland origin, with intense enhancement with gadolinium that had no intracranial involvement (fig. 1). The physical examination revealed a painless, nonpulsatile mass that displaced and enlarged the left naso-oropharyngeal wall. During a previous tracheotomy, a lateral cervicotomy associated with a transparotid and transmandibular approach was performed. The postoperative evolution was satisfactory and uneventful. The only complication observed was a 2-month paresis of a marginal branch of the facial nerve. Seven months later, an MRI revealed complete tumoral resection.

The histopathologic result revealed a highly cellular tumor, mildly pleomorphic, with round and oval to spindle cells with no atypia, very tightly packed in an intimate relationship with a profuse vascular network located in a collagenous stroma. Large vessels had a characteristic 'staghorn' morphology. Immunohistochemistry showed strong positivity with vimentin and CD34 and was negative for cytokeratins, EMA, S100 and others (fig. 2). Vimentin is the only marker expressed consistently in HPC [3].

Discussion

HPC is a very rare type of neoplasm, and when located in the parapharyngeal space, it becomes even more uncommon. Only a few cases have been reported previously in the worldwide literature (table 1). There is an equal incidence in both sexes, and the appearance
peak occurs in the fifth and the sixth decade of life [2, 3]. Distant metastasis rates range from 18 to 69% [3]. Some studies have mentioned trauma, hypertension and long-term steroid use to be possible etiologies. In our case, they were dismissed [3].

The initial clinical suspicion may be given by symptoms due to the compression of surrounding structures or by its external appearance [4]. It has been reported that all masses

<table>
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<th>First author, year</th>
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<tr>
<td>Case 1 Fountoulakis [2], 2011</td>
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<td>Case 2 Fareed [3], 2012</td>
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<td>Case 3 Dimri [5], 2010</td>
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<td>Case 4 Vo [10], 2007</td>
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<td>Case 5 Gierrek*, 2000</td>
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<td>Case 6 Llorente [7], 1999</td>
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<td>Case 7 Robb [4], 1987</td>
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<td>Case 8 Wakisaka [8], 2009</td>
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<td>Case 9 Kairemo [discussed in 7], 1991</td>
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<td>Case 10 Ozdzinski*, 1995</td>
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<td>Case 11 Shaia [6], 2006</td>
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<td>Case 12 McIlrath [discussed in 7], 1963</td>
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<td>Case 13 McMaster [discussed in 7], 1975</td>
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<td>Case 14 Rosignoli [discussed in 7], 1993</td>
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* Taken from two abstracts in English of articles written in Polish, which report 2 cases of hemangiopericytomas of the parapharyngeal space; not included in the reference list.
bulging in the oropharyngeal area were larger than 3 cm, and 20% of them were found incidentally during routine physical examinations or while the patient was undergoing tests or follow-ups for other pathologies. The clinical behavior of HPC is unpredictable, and it is usually described as a painless enlarging mass [2, 4, 5]. Vascular signs may include telangiectasia, pulsation or audible bruit [4]. Hypoglycemia, hypertension, hypokalemia secondary to hyperreninemia and cardiac failure are some systemic effects that have also been described in previous studies [4].

The diagnosis relies on the histological examination. Findings such as uniform vascular spaces surrounded by densely packed cells and the characteristic reticulin 'staghorn' pattern are typical of HPC [4–6]. In the WHO classification of soft tissue tumors, benign and malignant HPC have been recognized [4]. Criteria of malignancy [4] are presence of necrosis, high cellular density, presence of pleomorphic or immature cells, hemorrhage foci, high mitotic activity (more than 4/10) and a large macroscopic mass (more than 5 cm; some studies 6.5 cm) [2, 3, 6]. In our case, it was considered a mass with low malignant potential.

CT scanning and MRI may provide useful preoperative information about tumor features, extension, relationship with surrounding structures and osseous erosions (table 2). The use of contrast ascertains the vascularity of the tumor. MRI is essential for scheduling the surgical plan and should be considered the 'gold standard' for the study of this tumor. Nevertheless, imaging is not specific and has no pathognomonic findings [2, 5, 7, 8]. Fine-needle-aspiration cytology or biopsy is not advised in this type of tumor due to the high risk of bleeding [7].

Table 2. Radiological features of the tumors in the parapharyngeal space

<table>
<thead>
<tr>
<th></th>
<th>Morphology</th>
<th>CT</th>
<th>MR:T1</th>
<th>MR:T2</th>
<th>MR:contrast-enhanced T1</th>
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<tbody>
<tr>
<td>Prestyloid parapharyngeal space</td>
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<td></td>
<td>Second branchial cleft cyst</td>
<td>Well defined</td>
<td>Uniloculated cystic lesion</td>
<td>Hypointense</td>
<td>Hyperintense</td>
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<td></td>
<td>Cystic lymphangioma</td>
<td>Well defined</td>
<td>Multiloculated cystic lesion</td>
<td>Hypointense</td>
<td>Hyperintense</td>
</tr>
<tr>
<td>Minor or ectopic salivary gland tumors</td>
<td>Well-defined spherical mass</td>
<td>Hyperdense well-defined lesion</td>
<td>Hypointense</td>
<td>Hyperintense</td>
<td>Usually homogeneous enhancement</td>
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<tr>
<td>Neurogenic tumors</td>
<td>Well-defined spherical mass</td>
<td>Hyperdense well-defined lesion</td>
<td>Hypointense</td>
<td>Hyperintense</td>
<td>Intense homogeneous enhancement</td>
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<td>Infiltrative masses: soft tissue sarcomas</td>
<td>Ill defined</td>
<td>Slightly ill-defined lesions</td>
<td>Hypointense involving several adjacent spaces</td>
<td>Hypointense ill-defined lesion</td>
<td>Variable heterogeneous enhancement</td>
</tr>
</tbody>
</table>

Postyloid parapharyngeal space

| Neurogenic tumors | Well-defined spherical mass | Hyperdense well-defined lesion | Hypointense    | Hyperintense           | Intense homogeneous enhancement |
| Paragangliomas    | Well-defined oval mass      | Hyperdense well-defined lesion | Hypointense    | Hyperintense           | Intense homogeneous enhancement |
Some differential diagnoses include glomus tumor, hemangiomas, histiocytomas, angiosarcoma, schwannoma and others [2, 3, 9]. It is important to clarify that in our patient, the initial suspicion was a pleomorphic adenoma of a minor salivary gland. For preoperative differential diagnosis, a conventional angiography may be considered as well as if embolization is part of the plan [2, 5, 7].

A radical excision is the treatment of choice [3]. Radiotherapy should be considered in recurrent tumors, cases of incompletely excised lesions and tumors with malignant features [2, 5]. Chemotherapy has its basic role in distant metastatic disease. If a complete resection is made, there is a 100% median survival rate at 60 months [3, 10].

Conclusion

The accurate management and prognosis of this type of neoplasm are provided by the definite diagnosis obtained by a correct histopathologic assessment. The biological behavior of these tumors is quite peculiar, for there have been clinical reports of metastasis found even in benign, nonmitotic hemangiopericytomas.

A thorough imaging evaluation must be performed prior to surgery. Large tumors with significant vascularity should be evaluated with angiography, and preoperative embolization may be taken into consideration. An attempt for a surgical radical resection must always be intended. Long-term follow-up is very important due to its imprecise nature and previously reported late recurrences.

References