Juvenile nasopharyngeal angiofibroma

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DESCRIPTION

A 10-year-old boy visited the outpatient department of radiodiagnosis of our institution with complaints of progressive unilateral nasal obstruction, nasal discharge for 3 months and difficulty breathing while sleeping. There was history of epistaxis with 30–40 mL of blood loss and intermittent headaches relieved with analgesics. The patient was physically well-built with no clinical pallor; however, he breathed through his mouth. An external, non-tender facial deformity was noted as swelling on the right cheek (figure 1A). Endoscopic examination revealed a mass filling most of the right middle and inferior meatus extending up to the choana, showing active serous discharge and tinges of blood clots. The mass was soft and compressible, arising from the right lateral wall. On CT, a soft tissue density lesion was noted in the nasopharynx, right sphenoid sinus and ethmoid air cells, with lateral extension into the right infratemporal fossa through the right sphenopalatine fossa, causing its widening. There was an anterior bulging of the posterior wall of the right maxillary sinus with a decrease in its size (figure 1B). CT scan helps in determining invasion of the cancellous sphenoid bone, tumour staging and treatment planning. Soft tissue extension was determined on contrast-enhanced MRI with a large ill-defined, vividly enhancing lesion arising from the right sphenopalatine foramen, causing its widening, with extension into the right infratemporal fossa and masseter space (figure 2A). The lesion was heterogeneously hyperintense on T2/Fluid-attenuated inversion recovery (FLAIR) with a small cystic area and few hypointense foci within, giving a salt and pepper appearance (figure 2B), hypointense on T1, no restriction diffusion on Diffusion weighted imaging (DWI), and a few focal areas of blooming on Gradient-echo sequences (GRE).

The lesion extended into the posterior nasal cavity and ethmoid sinus, causing erosion of the right nasal cavity turbinate. Anterolaterally, anterior bowing of the posterior wall of the right maxillary sinus was noted, giving the pathognomonic Holman-Miller antral sign (figure 2B). The lesion extended into the right maxillary sinus and infratemporal fossa, causing widening of the pterygomaxillary fissure, with widening of the inferior orbital fissure and abutment of the right optic nerve apex. The superior and inferior extensions were up to the sella and hard palate, respectively, and the clivus posteriorly causing the supracavernous right internal carotid encaement. There was no infiltration into the buccal muscles. After imaging, the patient was referred for superselective embolisation.

Angiofibromas are benign, non-encapsulated lesions occurring in the nasopharynx, predominantly originating near the sphenopalatine foramen. They tend to extend in all directions with multiple projections. Epistaxis, unilateral nasal obstruction, pain and facial deformities are the most common symptoms associated with nasopharyngeal angiofibroma. They show connective tissue stroma with multiple vascular channels on histopathology. Surgery is required for the management of nasopharyngeal angiofibroma. Endoscopic nasal microsurgery is associated with less postoperative morbidity and blood loss during surgery but is restricted to limited lesions. Large tumours require transpalatal approach with lateral rhinotomy, and preoperative embolisation is performed to reduce intraoperative blood loss. Although large tumours require open surgery,
Images in...

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<th>Table 1</th>
<th>UPMC staging system for angiofibroma</th>
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<td>Stage</td>
<td>UPMC staging system</td>
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<tr>
<td>I</td>
<td>Nasal cavity, medial pterygopalatine fossa.</td>
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<tr>
<td>II</td>
<td>Paranasal sinuses, lateral pterygopalatine fossa; no residual vascularity.</td>
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<td>III</td>
<td>Skull base erosion, orbit, infratemporal fossa; no residual vascularity.</td>
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<tr>
<td>IV</td>
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<td>V</td>
<td>Intracranial extension, residual vascularity; M, medial extension; L, lateral extension.</td>
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UPMC, University of Pittsburgh Medical Center

Patient's perspective

I noticed a gradually increasing swelling on the right cheek of my son and he had difficulty in breathing with some foreign body sensation in his nose. We came to the rural hospital for further investigation and management. He was advised magnetic resonance imaging of PNS and orbit and diagnosis of juvenile nasopharyngeal angiofibroma was made. We are referred to a higher centre for further management.

Learning points

► Clinicians should not be deceived by the size of the tumour as perceived as it can only be the ‘tip of the iceberg’.
► Contrast CT aids in diagnosis and allows accurate staging of tumours, which is necessary in order to choose surgical techniques and estimate the prognosis.

Recent literature suggests that endoscopic endonasal removal can be performed postembolisation, for which the University of Pittsburgh Medical Center staging system is used (table 1).3

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Case reports provide a valuable learning resource for the scientific community and can indicate areas of interest for future research. They should not be used in isolation to guide treatment choices or public health policy.

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REFERENCES