Inflammatory Pseudotumor of the Nasopharynx

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CASE SUMMARY

An 87-year-old former smoker presented with severe unrelenting headache following trivial trauma. The patient also had developed recent onset facial numbness and facial palsy. Medical history included hypertension and hypothyroidism, with occasional alcohol ingestion. On examination, patient was cachectic with evidence of muscle wasting but examination was otherwise normal. There was no lymph node enlargement. Nasal examination was suspicious for a left nasopharyngeal mass, which was confirmed by nasopharyngeal endoscopy. Two biopsies of the nasopharyngeal mass demonstrated only fibrous tissue and mild chronic inflammation with no evidence of malignancy.

IMAGING FINDINGS

Nasopharyngeal inflammatory pseudotumor (IPT) appears as homogeneously enhancing nasopharyngeal soft tissue on contrast-enhanced CT examination. Associated destruction of the skull base (Figure 1) and narrowing of the neighboring skull base foramina/canals may occur. There is usually no enlargement of the cervical lymph nodes.

On MRI, IPTs are typically ill-defined, infiltrative masses (Figure 2) with moderate, homogeneous contrast enhancement. There is involvement of the submucosa of the nasopharynx with preservation of normal hyperintense signal of the nasopharyngeal mucosa. Contiguous structures that



FIGURE 1. (A) Axial contrast-enhanced CT scan of the nasopharynx demonstrates enhancing nasopharyngeal soft tissue (arrow) predominantly involving the left side. (B) Axial CT bone window shows destruction of the clivus (arrow), extending across the midline.

may be involved include the clivus, eustachian tube, carotid and parapharyngeal spaces, Meckel cave, orbital apex, and cavernous sinus. Encasement of the internal carotid artery may occur. Ipsilateral opacification of the mastoid air cells is a common finding due to eustachian tube obstruction.

DIAGNOSIS

Inflammatory pseudotumor (IPT) of the nasopharynx.

Differential diagnosis includes nasopharyngeal carcinoma (NPC), chordoma, chondrosarcoma, skull base lymphoma, plasmacytoma, and metastasis. Rosai-Dorfman disease, Epstein-Barr virus-related inflammatory pseudotumor, calcifying fibrous pseudotumor, and sclerosing lymphoma are less likely considerations.

DISCUSSION

Inflammatory pseudotumor (IPT) of the nasopharynx is a benign, idiopathic disease usually mistaken for a neoplasm/infection owing to its aggressive appearance^{1,2,3} Patients routinely present with recent history of otitis media, hearing loss, and/or single or multiple cranial nerve neuropathies.²

IPT is an inflammatory lesion of unknown etiology. Various theories, including autoimmune reaction linked to viral infection, infectious process due to sinusitis, or syphilis and aberrant production of fibrogenic cytokines, have been proposed.¹ However, recent studies have suggested cytogenetic clonal abnormality and anaplastic lymphoma kinase expression, suspicious for a neoplastic cause.⁴



FIGURE 2. (A) Axial T1 MRI through the nasopharynx shows a soft-tissue mass lesion of the left posterolateral aspect of the nasopharynx (arrow) with destruction of the clivus and left petrous apex. (B) Axial T2 MRI of the nasopharynx demonstrates the low signal in the lesion (arrow). (C) The left nasopharyngeal mucosa (arrow) is unremarkable on axial T2 MRI of the nasopharynx, favoring IPT over NPC.



FIGURE 3. Hematoxylin and eosin stains on microscopy show (A) Neutrophils with areas of necrosis (smudgy purple areas) and fibrous tissue (pink). (B) Lymphocytes (small blue round dark blue cells) are intermingled with plasma cells (slightly larger nuclei with larger pink cytoplasm).

On gross pathology, IPTs resemble inflamed fibrous tissue with contiguous reactive bone. Routine microscopy demonstrates mixed acute/chronic inflammatory cells and spindle cells with a densely fibrotic background.^{5,6} (Figure 3).

On MRI, IPT is typically seen as a mass which is T2 hypointense compared to brain parenchyma (Figure 2), while NPC (the most common differential diagnostic consideration) is more hyperintense on T2 imaging.^{1,4} Internal carotid artery encasement and narrowing of the vessel lumen are common features with IPT.¹ In comparison to NPC, IPT involves the submucosal layer of the nasopharynx. The hyperintense mucosal layer is well-delineated from the hypointense submucosal lesion on T2 imaging (Figure 2). Extensive pachymeningeal thickening and enhancement is a more common feature of IPT but not of NPC. Cervical lymphadenopathy is more commonly encountered in patients with NPC than with IPT.^{1,2,6} Systemic steroids are the treatment of choice in patients with IPT involving the nasopharynx and skull base.⁷

CONCLUSION

Inflammatory pseudotumor of the nasopharynx is a rare mimicker of nasopharyngeal carcinoma and other neoplastic and infectious lesions involving the skull base.

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