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A rare mass in the nasopharynx: Metastatic small cell lung cancer masquerading as nasopharyngeal fullness

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Abstract

Small Cell Lung Cancer (SCLC) is notorious for its aggressively proliferative nature. Although early metastasis is welldocumented in SCLC, metastasis to the nasopharynx is exceedingly rare.

We present the case of a 63-year-old male with a history of SCLC of the lung treated with Chemo Radiation Therapy (CRT) at the time of diagnosis who presents 2 years later with rapidly worsening weight loss, dyspnea, dysphagia, dysphonia, right-sided V3 neuralgia, and right-sided nasal obstruction. Prior to presentation, the patient had poor compliance with CRT and was lost to follow-up. Endoscopy showed a hyperemic mass encompassing the entire posterior nasopharynx and the sphenoethmoidal recess with right true vocal cord paralysis. Imaging revealed an advanced nasopharyngeal mass with extension to the deep lobe of the right parotid gland and complete effacement of the parapharyngeal space. Biopsy of the mass revealed malignant epithelial cells arranged in sheets with cells staining positive for synaptophysin, confirming neuroendocrine differentiation and likely metastasis from incompletely treated SCLC. Due to the extent of disease, multi-disciplinary recommendations were made to begin CRT.

The literature regarding nasopharyngeal metastasis of SCLC is scarce. Our case serves to highlight this rare anomaly and demonstrate the significance of patient compliance in accomplishing successful therapeutic management.

Introduction

Small Cell Lung Cancer (SCLC) is strongly associated with exposure to tobacco products and is marked by an exceptionally high proliferative rate, strong predilection for early metastasis, and poor prognosis [1]. Common routes of metastasis of SCLC include lymphatic spread to hilar and mediastinal lymph nodes and hematogenous spread to the contralateral lung, brain, liver, adrenal glands, and bone [2]. While SCLC metastasis to distant organs is well-documented, occurrences of metastasis to the nasopharynx are exceedingly uncommon and warrant further exploration.

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Understanding the underlying mechanisms and clinical implications of rare metastatic disease is imperative to enhance diagnostic precision and tailor effective treatment strategies [3]. Early chemo radiation is the standard of care for limited-stage SCLC. Although most SCLCs initially respond to therapy, they have a propensity to recur [4]. This case furthermore exemplifies the critical role of early chemo radiation therapy in managing metastatic SCLC, emphasizing the importance of patient compliance in achieving optimal therapeutic outcomes [5-7]. In this case report, we aim to shed light on a rare SCLC metastasis and demonstrate the significance of patient engagement and support to achieve successful therapeutic interventions.

Case presentation

In this case report, we present a 63-year-old SCLC patient who was treated with chemo radiation to the lung at the time of initial diagnosis. However, he had a history of poor compliance with numerous missed treatments and was ultimately lost to follow-up. He subsequently presented two years later with rapidly worsening unexpected weight loss, dyspnea, dysphagia, dysphonia, right-sided V3 neuralgia, and right-sided nasal obstruction.

Flexible nasal endoscopy was significant for a hyperemic mass encompassing the entire posterior nasopharynx and sphenoethmoidal recess (Figure 1). Evaluation of the right true vocal cord showed paralysis and fixation in the paramedian position with a mobile left true vocal cord. Functional endoscopic evaluation of swallowing revealed significant residual pooling in the piriform sinuses and vallecula with penetration and aspiration of liquids. CT imaging revealed a locoregionally advanced nasopharyngeal mass with lateral extension to the deep lobe of the right parotid gland and complete effacement of the parapharyngeal space. Retropharyngeal lymphadenopathy was noted without any lateral cervical lymph nodes (Figure 2). The patient was admitted for nutritional optimization and a gastrostomy tube was placed during his inpatient stay.

Biopsy of the right nasopharyngeal mass revealed malignant epithelial cells arranged in sheets. The tumor cells were cohesive with high nuclear-to-cytoplasmic ratios, hyperchromatic nuclei, and granular chromatin. The cells were positive for synaptophysin confirming neuroendocrine differentiation (Figure 3). After discussion at a multidisciplinary tumor conference and confirmation with the pathology team, consensus was reached that this was likely metastatic disease from incompletely treated SCLC.

Due to the extent of metastatic disease, multi-disciplinary recommendations were made for referral back to the oncology team for consideration of systemic therapy, the radiation oncology team for possible palliative radiation to the nasopharynx, and supplemental imaging studies to rule out intracranial involvement. Since the patient had concurrent moderate-to-severe conductive and sensorineural hearing loss, he was deemed to not be a good candidate for cisplatin/etoposide. Thus, carboplatin and etoposide chemotherapy was initiated along with radiation therapy. Magnetic resonance imaging two months after the CT scan revealed the nasopharyngeal mass, showed a progression of the solid mass arising from the right nasopharynx. The mass measured approximately 7.1 x 9.3 x 4.5 cm and was shown to obliterate the right parapharyngeal space, invade the adjacent right medial pterygoid muscle, encase the right internal carotid artery, obliterate the internal jugular vein, and invade the right side of the clivus. The mass was still shown to invade the deep and part of the superficial lobes of the right parotid gland (Figure 4). The mass also infiltrates the longus capitis muscles, right greater than left, and extends into the right nasal cavity and right pterygopalatine fossa. The previously noted right lateral retropharyngeal metastatic lymph node conglomerate was demonstrated on MRI to be congruent with the nasopharyngeal mass. No discrete intracranial extension is noted. No orbital or hypopharyngeal involvement is seen.

Since the initiation of chemoradiation therapy, the patient's medical course has been complicated by hospital admission due to neutropenic infection (low WBC of 0.44). Multi-disciplinary medical treatment with oncology, radiation oncology, otolaryngology, and pulmonology continues as therapeutic management is maximized.



Figure 1: Endoscopic view of nasopharyngeal mass through the left nasal cavity.

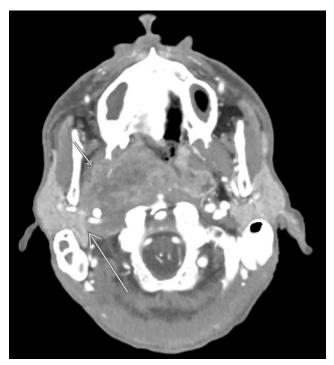


Figure 2: CT imaging demonstrating nasopharyngeal mass extending to deep lobe of right parotid gland.

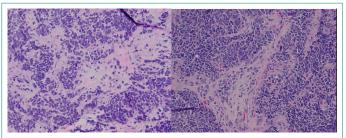
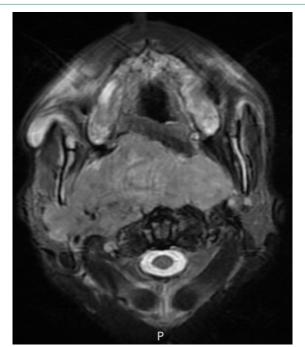
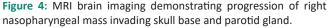


Figure 3: Side-by-side pathology slides demonstrating neuroendocrine cell populations from nasopharyngeal biopsy and previous small cell lung cancer.





Discussion

Small Cell Lung Cancer (SCLC) is renowned for its aggressive nature, rapid proliferation, and propensity for early metastasis, often before clinical detection. Most patient have metastasis at the time of diagnosis, with less than 50% two-year survival and less that 5% five-year survival [8]. The high mitotic rate of SCLC lends to its aggressive behavior, contributing to the early formation of distant metastases [1]. The feature of propensity to metastasize sets SCLC apart from Non-Small Cell Lung Cancer (NSCLC) [9]. The most common sites of metastasis for SCLC include the liver, bones, adrenal glands, and brain [2]. The tumor's extensive neuroendocrine features and high growth fraction contribute to the hematogenous and lymphatic spread [10].

Metastasis of SCLC to the nasopharynx represents a rare and intriguing clinical picture, diverging from the more typical sites of spread. The exact mechanism underlying this uncommon metastasis remains unclear, but one possible route could be hematogenous dissemination due to the rich vascularity and anatomical position of the nasopharyngeal region, allowing circulating tumor cells to infiltrate and establish metastatic lesions [11].

Management of SCLC requires comprehensive therapeutic strategies [3]. For patients with limited-stage SCLC, the standard of care involves early initiation of chemoradiation therapy. Combining chemotherapy, often platinum-based, with radiation aims to eradicate both local and systemic disease [7,12]. The success of this treatment modality is evident in the initial response of most SCLCs to therapy. However, despite a remarkable early response to treatment, recurrence is common [4].

The prominent trait of recurrence in SCLC draws attention to the need for consistent follow-up and vigilance in monitoring patients during and after treatment. Early detection of recurrence is crucial for ensuring swift therapy, potentially improving outcomes and quality of life for patients [5]. Adherence to recommended follow-up appointments, surveillance imaging, and, most importantly, chemo-radiation appointments is essential for optimizing outcomes [13]. Because SCLC requires a proactive treatment plan, a multi-disciplinary approach and efficient communication between providers and patients are vital for patient compliance and successful long-term management [6].

Conclusion

This case provides a prime example of the importance of consistent follow-up and compliance and some of the devastating, albeit rare, effects that can result from incompletely treated SCLC.

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