doi:10.4317/medoral.21506 http://dx.doi.org/doi:10.4317/medoral.21506

Orbital apex syndrome affecting head and neck cancer patients: A case series

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Received: 28/06/2016 Accepted: 15/02/2017 Prado-Ribeiro AC, Luiz AC, Montezuma MA, Mak MP, Santos-Silva AR, Brandão TB. Orbital apex syndrome affecting head and neck cancer patients: A case series. Med Oral Patol Oral Cir Bucal. 2017 May 1;22 (3):e354-8.

http://www.medicinaoral.com/medoralfree01/v22i3/medoralv22i3p354.pdf

Article Number: 21506 http://www.medicinaoral.com/ © Medicina Oral S. L. C.I.F. B 96689336 - pISSN 1698-4447 - eISSN: 1698-6946 eMail: medicina@medicinaoral.com Indexed in: Science Citation Index Expanded Journal Citation Reports Index Medicus, MEDLINE, PubMed Scopus, Embase and Emcare Indice Médico Español

Abstract

Background: Orbital apex syndrome (OAS) is a complex and uncommon disorder that typically damages multiple cranial nerves in association with optic nerve dysfunction. OAS is associated with several different pathologies, however; only a few cases have been reported in association with head and neck cancer (HNC) so far.

Material and Methods: A case series of HNC patients diagnosed with OAS is described including clinicopathological data, image findings, and disease outcome.

Results: Ptosis and diplopia were diagnosed in four male patients with mean age of 61.2 years who were undergoing treatment for late-stage carcinomas of the tongue, larynx, and nasopharynx, eventually leading to the diagnosis of OAS. The mean overall survival rate after the diagnosis of OAS was 9.5 months.

Conclusions: The current study reinforces evidence that OAS indicates poor prognosis and highlights the importance of early diagnosis.

Key words: Head and neck cancer, oral cancer, metastasis, orbital apex syndrome, optic neuropathy.

Introduction

Orbital apex syndrome (OAS) is a rare disorder characterized by the involvement and damage of cranial nerves, including the oculomotor (III), trochlear (IV), abducens (VI) and ophthalmic branch of the trigeminal nerve (V) in association with optic dysfunction (1-3). OAS may be associated with inflammatory diseases (sarcoidosis, systemic lupus, Wegener granulomatosis), fungal infections (*Mucormycosis, Aspergillus*), bacterial infections (*Actinomyces, Mycobacterium tuberculosis*), viral infections (*Herpes zoster*), primary and metastatic tumors (nasopharyngeal carcinoma, neural

tumors, metastatic tumors, lymphomas and leukemia) as well as trauma (sinonasal surgery), among others (mucoceles) (1,2).

From a clinical point of view, patients diagnosed with OAS initially present visual loss and ophthalmoplegia, such as proptosis, ptosis, fixed dilated pupil, and diplopia. Associated periorbital pain may also develop in these patients (1-3).

Apparently, the association of OAS and head and neck cancer (HNC) is considered to be exceptionally rare, with only a few cases published in the English-related literature so far (4,5). Therefore, the aim of the present case series was to report four additional cases of OAS diagnosed in patients undergoing treatment for advanced carcinomas of the head and neck region, including tumors of the tongue, larynx, and nasopharyns. In addition, clinic and image data relevant for diagnosis will be discussed in the light of the scientific literature pertinent to OAS.

Material and Methods

This is a retrospective case description report based on patients who were undergoing dental treatment at the Dental Oncology Service of the Instituto do Cancer do Estado de Sao Paulo (ICESP), from March 2011 to January 2013. In order to be included in this case series, the patients had to present a confirmed diagnosis of HNC and develop ocular alterations or ophthalmoplegia. Complete medical information including demographic and clinicopathologic data, diagnostic head and neck imaging and post-cancer treatment follow-up also had to be complete and fully available at the patients' medical charts.

This retrospective study was reviewed and approved by the Ethics Committee of the University of Sao Paulo Medical School, Sao Paulo, Brazil (study protocol number 882,731).

Results

Four patients with a previous history of head and neck carcinomas were included in this case series. All of the patients were men, with a mean age of 61.2 years at the time of diagnosis, presenting advanced stage tumors with a previous history of tobacco and alcohol consumption. Detailed demographic and clinicopathological information are presented in tables 1 and 2. Illustrative images of the patients are presented in figures 1,2 and 3.

Discussion

HNC is the sixth most common cancer worldwide (6,7), accounting for an annual incidence of approximately 700,000 cases (nearly 5% of all cancers) and almost 375,000 deaths (7). HNC primarily affects men between the sixth and the seventh decades of life. Smoking and alcohol use are the major risk factors (8). In accordance

Table 1 Demographic and cliniconathologica	l features of head and neck cancer	patients diagnosed with orbital apex syndrome.
rable 1. Demographic and enneopathologica	reatures of neau and neek cancer	patients diagnosed with orbital apex syndrome.

Patient	Gender, age	Cancer diagnosis	Clinical Stage	Cancer treatment	Cancer risk factors
1	Male, 73 years old	Tongue (lateral border) squamous cell carcinoma	Т3N2bM0	Partial Glossectomy and radical bilateral neck dissection; 3D conformal radiothera- py (66 Gy); chemother- apy (Carboplatin)	Tobacco (20 cigarettes/ day, during 57 years) and alcohol (1 L distilled spir- its/day, during 50 years) consumption
2	Male, 70 years old	Larynx squamous cell carcinoma	T4N3M0	3D conformal radio- therapy (70 Gy)	Tobacco (40 cigarettes/ day, during 10 years) and alcohol (1.5 L distilled spirits/day, during 50 years) consumption
3	Male, 53 years old	Nasopharyngeal carcinoma	T4N3M1 (liver metastasis)	Palliative treatment: 3D conformal radio- therapy (40 Gy); che- motherapy (Cisplatin and 5 fluorouracil)	Tobacco (20 cigarettes/ day, during 12 years) and alcohol (1 L distilled spir- its/day, during 12 years) consumption
4	Male, 49 years old	Nasopharyngeal carcinoma	T4N3M1 (bone metastasis)	Palliative treatment: 3D conformal radio- therapy (30 Gy); che- motherapy (Cisplatin and 5-fluorouracil)	Tobacco (20 cigarettes/ day during 30 years) smoking

Patient	Diagnosis timing	Ocular alteration	Image features	Involved Cranial Nerve	Medical follow-up
1	4 months after chemoradio- therapy	Diplopia; facial paresthesia (right side, ipsilateral to tumor) {Figure 1A}	Computed tomography: hyperdense image localized in the base of the skull associated with destruction of the right sphenoid bone (Figure 1B)	Abducens (VI)	Disease progression with progressive ptosis and sensory ataxia (denied pain); died 9 months after the diagnosis of OAS
2	During radiotherapy (19 th session, 38 Gy)	Ptosis; ophthalmoplegia and hearing loss (left side, ipsilateral to tumor) {Figure 2A}	Computed tomography: thickening of the left pinna and external conduit (Figure 2B)	Oculomotor (III); Vestibulocochlear (VIII)	Died 9 months after the diagnosis of OAS
3	During radiotherapy (2 nd session, 5 Gy)	Ptosis; blurred vision; mydriasis and ophthalmoplegia (left side, ipsilateral to tumor) {Figure 3A}	Magnetic resonance imaging: hyperintense image at the skull base causing enlargement of the left optical canal and destruction of the left sphenoid bone (Figure 3B)	Oculomotor (III)	Died 9 months after the diagnosis of OAS
4	Concomitant to cancer diagnosis	Diplopia; progressive visual loss (left side, ipsilateral to tumor) {Figure 3C}	Magnetic resonance imaging: mass involving the left sphenoid bone and causing hypoglossal canal obliteration (Figure 3D)	Trochlear (IV); Hypoglossal (XII)	Died 11 months after the diagnosis of OAS

Table 2. Clinicopathological features of orbital apex syndrome in head and neck cancer patients.

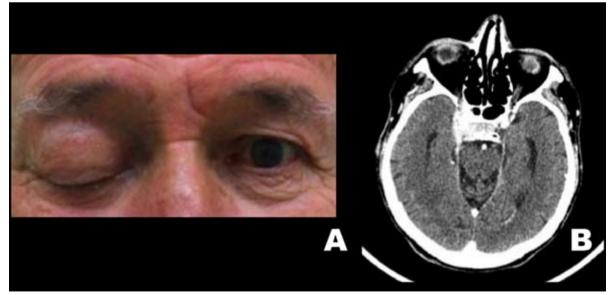


Fig. 1. A. Extraoral clinical aspect showing right eyelid ptosis. B. Axial CT (soft-tissue window display) demonstrating an extensive hyperdense image at the base of the skull associated with the sphenoid bone.

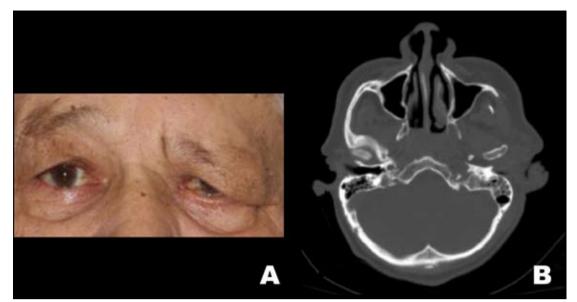


Fig. 2. A. Extraoral clinical image of the left eyelid ptosis. B. Axial CT (bone-window display) showing a thickening of the left pinna and external ear conduit.

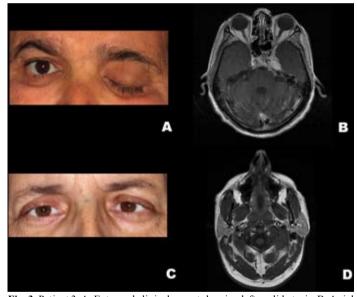


Fig. 3. Patient 3. A. Extraoral clinical aspect showing left eyelid ptosis. B. Axial IMR (T2-weighted) demonstrating hyperintense image at the skull base causing enlargement of the left optical canal. Patient 4. C. Extraoral clinical aspect revealing fixed globe in the left eye. D. Axial IMR (T2-weighted) demonstrating a mass involving the left sphenoid bone and causing hypoglossal canal obliteration.

with the above-mentioned literature, all the patients in the current series were elderly men with a previous history of tobacco and alcohol abuse.

HNC patients tend to present a loco-regional spread of disease; the most common site of metastasis is the regional neck lymph nodes. Distant metastasis from HNC is considered a rare event (9-11). Kowalski *et al.* (10) evaluated 2,327 patients diagnosed with HNC and found

that only 3.8% of the patients had distant metastasis, while Alvi *et al.*, (9) reported a distant metastasis rate of 23%. Lung and bone were the most common distant sites for distant metastasis in both studies. Accordingly, advanced HNCs affected all of the patients included in the present case series.

The association between OAS and HNC is unusual. In this context, Whirth *et al.*, (5) recently reported a case

of oral squamous cell carcinoma associated with ophthalmoplegia and severe general facial pain, leading to the diagnosis of OAS. Aryasit *et al.* (2) performed a retrospective review of patients diagnosed with OAS and observed that 48% of which had associated neoplasias were lymphomas (37.5%) was the most common tumor followed by meningiomas (29.2%). Apparently, the incidence of the association between HNC and OAS is currently unknown and, when judging the cases presented herein, one could suggest that this association might be unrecognized and might not be as rare as once believed.

Patients diagnosed with OAS usually present with visual impairment and ophthalmoplegia, including proptosis, ptosis and periorbital pain. The diagnosis of OAS is clinical; however, a detailed history and physical examination should be performed (12,13). Imaging is important in discerning the causative etiology (2,3). In the current case series, all the patients presented ocular alterations, including two patients presenting diplopia and two other presenting ptoses.

Most of the patients diagnosed with distant metastasis of HNC are considered to be incurable and are often treated by palliative care protocols, such as chemotherapy or radiotherapy (10). According to Alvi *et al.* (9), the average overall survival rate after the diagnosis of distant metastasis of HNC is 5 months, while Kowalski *et al.* (10) reported a rate of 1.8 months in patients with bone metastasis. In the present case series, the average overall survival rate after the diagnosis of skull metastasis (or infiltration) associated with OAS was 9.5 months. The present results reinforce previous evidence that OAS is a marker of poor disease outcome in cancer patients and emphasize the need for prompt identification of clinical signals of OAS in HNC patients (2).

Overall, up to 80% of all HNC patients are diagnosed with advanced stages of the disease, presenting poor prognoses and requiring aggressive multimodality treatment (7,12,14) in association with multidisciplinary supportive care where dental oncologists may contribute to the early diagnosis of the disease relapse or relevant complications of tumor progression, such as OAS. Thus, this case series highlights the importance of interactions among dentists, oncologists, and physicians involved in HNC treatment to improve the early recognition of OAS and promote patient prognosis.

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Acknowledgements

Supported by Fundação de Amparo à Pesquisa do Estado de São Paulo (FAPESP, processes numbers 2012/06138-1 and 2013/18402-8).

Conflict of Interest

The authors have declared that no conflict of interest exist.