

Giant nasopharyngeal tumor revealed by unilateral sixth nerve palsy : A case report

Abstract :

Nasopharyngeal tumor is an intrusive tumor arising from the epithelium of the nasopharynx, presenting as different symptoms depending of the area of the development of the tumor.

The purpose is to report a rare neuro-ophthalmic presentation of nasopharyngeal carcinoma .

We report the case of a 60 years old woman who first presented with a sixth nerve palsy leading to the diagnosis of a giant nasopharyngeal tumor extending to the cavernous sinus.

Nasopharyngeal tumor is an endemic carcinoma in North Africa. The location of the tumor, its difficult access to the physical examination and the symptomatology often misleading associated with invasion of adjacent structures explain the often delayed diagnosis.

Keywords : Nasopharyngeal tumor – Abducens – Diplopia – cavernous sinus

Introduction :

Nasopharyngeal carcinoma is one of the most frequent cancers in North Africa. The most common histological type is non-keratinising squamous cell carcinoma.

The intracranial extension may be revealed by cranial nerve paralysis or orbital apex syndrome . Cranial nerve involvement isn't rare but the isolated sixth nerve paralysis as first presentation of the tumor is uncommon.

Case report :

We report the case of a 70 years old female , with no history of diabetes or cardiovascular disease who presented with a 4-week history of limited eye movement plus diplopia .

The ophtalmological examination found limitation in abduction of the left eye and an horizontal binocular diplopia worsened with left gaze . (Fig1)

The best-corrected visual acuity (BCVA) was 20/25 in both eyes, pupil was reactive and equal . Exam of the anterior and posterior segment were unremarkable .



Fig 1 : Patient examination showed significant abduction deficit of the left eye

Orbitocerebral magnetic resonance imaging (coronal and axial slices) was performed and showed a giant left nasopharyngeal tumor starting to extend to the left cavernous sinus. (Fig2)

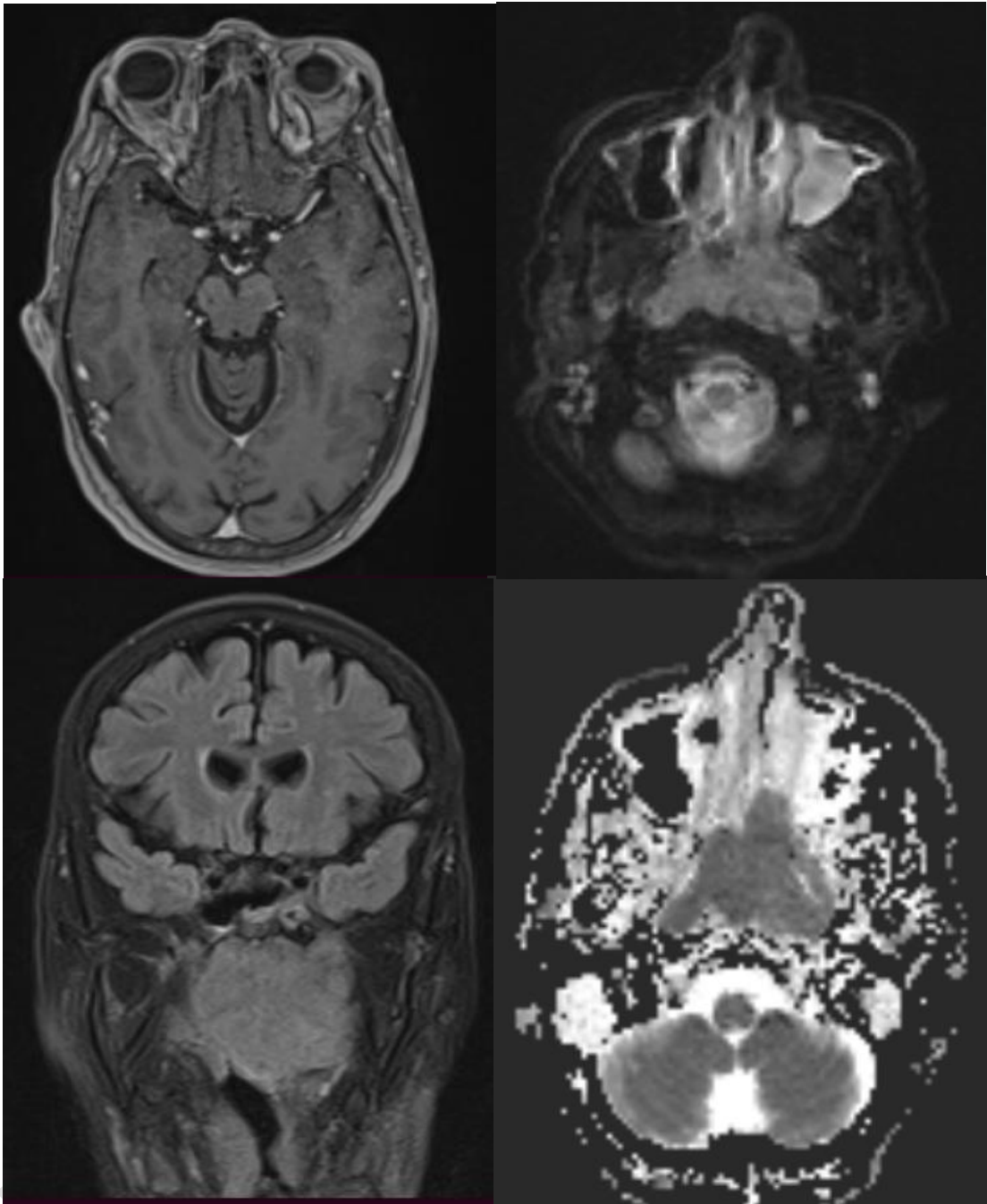


Fig 2 : Giant nasopharyngeal tumor : Isosignal on T1 sequences , hypersignal on diffusion sequences with low ADC with significant enhancement after gadolinium injection .

The patient was referred to an otolaryngologist who performed a rhinoscopy that showed a mass at the left nasopharynx .Tumor biopsy was done and revealed a non keratinizing squamous cell carcinoma .(Fig 3)

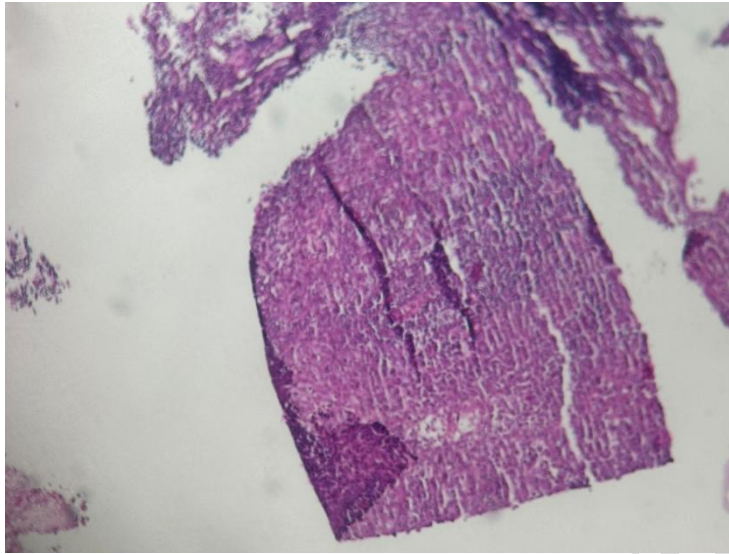


Figure 3 : Histopathological examination revealing a non keratinizing squamous cell carcinoma

Discussion :

Nasopharyngeal tumors are frequent tumors in North Africa with an incidence rate of 8-12/10000. The tumor occurs in the epithelial lining of the nasopharynx .On a pathological basis , there are 3 histological types and the undifferentiated nonkeratinizing NPC is the most common type . [1]

Almost half of the clinical presentations of the nasopharyngeal tumor are nasal symptoms although other symptoms may be present depending on the extension . (Table1)

Structures	Clinical presentation
Nasal	Nasal obstruction/Epistaxis
Aural	-Hearing loss -Tinnitus
Neurological	-Diplopia (Sixth nerve) -Opthalmoplegia (Cranial nerves) -Horner's syndrome -Xerophthalmia [15]
Metastasis	-Cervical lymph node - Paraneoplastic syndrome

Table 1 : Different clinical symptoms of nasopharyngeal carcinoma[2]

Cranial nerve palsy represents 9.4% symptoms of NPC.[2]. Cranial neuropathy is usually associated with a more invasive stage T4 [3] which explains the poor prognosis in cranial nerve involvement. The cranial nerves that are the most involved are nerve V (particularly maxillary division) and VI [4]. A study conducted by Richards et al that found NPC was responsible for abducens palsy in (5.6%) out of 193 patients with ocular motor nerve palsy caused by intracranial neoplasms.

In most cases in literature, abducens palsy associated with diplopia is only present as a sign of the recurrence of the tumor or post radiotherapy. Kau and al analysed a series of patients presenting diplopia secondary to the tumor. They found that new onset diplopia in these patients was secondary to tumor recurrence in (52%) of the patients, radiation neuropathy in (35%), and skull base osteoradionecrosis in (13%).[5]

Wong and al analysed data of 354 patients diagnosed with or treated for NPC. 7.6% of the patients presented an ophthalmic involvement due to NPC or its treatment. Symptomatic orbital invasion occurred in 48.1%. Ocular signs, but no orbital invasion, were present 29.6%. [6]. However cavernous sinus infiltration was not statistically significant as a factor of poor prognosis in patients treated for NPC even in advanced lesions [7].

CT scan supplemented with MRI is currently being used for accurate tumor mapping and detection of possible tumor extension [7].

In a series of 166 patients, Ilhan and al tried to analyse the outcome of abducens nerve paralysis in patients with nasopharyngeal carcinoma. The third nerve palsy was associated with poor outcome compared to the sixth nerve palsy that is more amenable to treatment. [8] The duration of neuropathy and the involvement of the optic nerve were independent negative predictors for neurological recovery because of the irreversible damage to the neural tissue, even if the underlying cause is addressed. [9]

Nasopharyngeal tumors are not amenable for surgery because of the anatomic specificity of the region, but it is a highly radiosensitive tumor. Radiotherapy or chemoradiotherapy is considered the first choice treatment for NPC.[10]

According to several studies, NPC misdiagnosis can be attributed to two causes: patient delay and health system issues (professional delay)[11]. Improper diagnosis and a lack of suspicion of NPC by healthcare professionals, particularly in low-incidence areas, contribute to late diagnosis.[12] The anatomical specificity of the tumor is also related to a late diagnosis. James and al: in a series of 504 patients the median time between symptoms and diagnosis of NPC was 2 months [15]

Conclusion :

Nasopharyngeal carcinoma is among the most frequently misdiagnosed tumors since it does not initially present with nasal symptoms. While early optic nerve involvement is extremely rare, symptoms are associated with cranial involvement. To improve patient survival, NPC should be diagnosed early and any isolated nerve palsy should lead to neuroimaging.

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