Sphenoid sinus organized hematoma with cranial neuropathies masquerading as a malignancy: A case report

YU-HSUAN LIN1, PO-CHIN WANG2 and YAOH-SHIANG LIN1,3

Departments of 1 Otolaryngology - Head and Neck Surgery, and 2 Radiology, Kaohsiung Veterans General Hospital, Kaohsiung 813; 3 Department of Otolaryngology - Head and Neck Surgery, National Defense Medical Center, Taipei 114, Taiwan, R.O.C.

Received November 19, 2014; Accepted October 5, 2015

DOI: 10.3892/ol.2016.4475

Abstract. Sinonasal organized hematoma (SNOH) is rarely encountered in clinical practice. The disease demonstrates a high tendency for occurrence in East Asian individuals, and in the majority of cases, is located in the maxillary sinus. The current report presents the case of an 81-year-old female who developed a space-occupying lesion, which masqueraded as a skull base malignancy, following surgery for the treatment of isolated sphenoid sinus aspergilloma. Subsequent endoscopic endonasal surgery confirmed the diagnosis of an OH of the sphenoid sinus. The patient recovered from all neurological deficits within two months, with the exception of the loss of visual perception. Although SNOH presents a diagnostic challenge, when physicians possess knowledge of its typical imaging features, this facilitates the achievement of a correct diagnosis and the prescription of optimal treatment.

Introduction

Sinoonasal organized hematoma (SNOH) was initially described in 1917 by Tadokoro (1) as a benign lesion characterized by an expansile nature and locally aggressive behavior, capable of mimicking a potentially dangerous neoplastic process (2-5). The diagnosis of SNOH is challenging, even given the integration of multiple informative symptoms (2). Computed tomography (CT) and magnetic resonance imaging (MRI) are useful for the assessment of potential cases of SNOH (2,5), and usually reveal typical bony changes and a distinct speckled pattern that exhibits various signal intensities, respectively. The hallmarks of SNOH are typically a dark peripheral rim on T2-weighted imaging (T2WI), and a nodular and patchy enhancement in post-contrast T1-weighted imaging (T1WI) (2,5). The successful treatment of SNOH relies on complete surgical excision (3,5). However, one notable and unexplained anomaly is that the disease demonstrates a high tendency for occurrence in the maxillary sinus (3,5-7). Only one case involving the sphenoid sinus, an intricately structured space in close vicinity to vital structures, has been reported to date (4). To the best of our knowledge, the current report presents the first case of an OH of the sphenoid sinus with multiple cranial nerve involvement, which was identified following surgery for the treatment of isolated sphenoid sinus aspergillosis (ISSA). The disease was successfully treated using endoscopic endonasal surgery, yielding a positive outcome for the patient.

Case report

An 81-year-old woman presented as an inpatient to the Kaohsiung Veterans General Hospital (Kaohsiung, Taiwan) in October 2013, with a 2-year history of an excruciating headache, which was described as deep-seated and throbbing. The patient had subsequently developed progressive deterioration of visual acuity in the right eye and drooping of the right eyelid, although the total duration of these symptoms was uncertain. The medical history of the patient noted the use of endoscopic sinus surgery for the treatment of ISSA 3 years previously, and a long-standing history of chronic kidney disease. Upon examination, besides optic atrophy, the patient's right eye exhibited complete ptosis and mydriasis with a sluggish pupillary reaction. The eye was positioned looking down and out, and the orbit could not be adducted. These symptoms were compatible with a diagnosis of compressive oculomotor nerve palsy. The nasal endoscopy revealed a friable mass with superficial telangiectasia straddling the torus tubarius and extending forward (Fig. 1). Non-contrast CT revealed an aggressively enlarging lesion in the midline skull base, with destructive bony structures (Fig. 2). Distinctive internal heterogeneous hyperintensity
and a hypointense peripheral rim on T2WI, and a patchy enhancement pattern on post-contrast T1W1 were observed (Fig. 3). Routine laboratory data revealed normal coagulation and pituitary function profiles, including human growth hormone (0.23 ng/ml; normal range, 0.0-16 ng/ml), luteinizing hormone (31.35 mIU/ml; normal range, 11.3-38.7 mIU/ml), follicle stimulating hormone (74.09 mIU/ml; normal range, 31.8-134 mIU/ml) and free thyroxine (1.58 ng/dl; normal range, 0.8-1.9 ng/dl). Only high sensitivity-thyroid-stimulating hormone levels were abnormal (0.245 uIU/ml; normal range, 0.4-4.0 uIU/ml).

In order to optimize surgical planning, the patient underwent a biopsy, which yielded a negative result. The patient subsequently underwent endoscopic endonasal surgery to resect the lesion; a posterior ethmoidectomy, sphenoidotomy and septoplasty were required in order to gain complete surgical access. A well-capsulated, russet-colored and blood clot-containing tumor was removed in a piece-by-piece manner, leaving the periosteal layer of dura mater that protected the intracranial structures intact. The thinned bony components located above the right orbital apex and right optic canal were gently flaked off. The surgical extent was explored inferiorly to the upper clivus. A sample of the lesion sent for pathological analysis revealed an OH without any evidence of fungal hyphae (Fig. 4). The immediate post-operative course was uncomplicated, with resolution of the patient's headache. At the 2-month follow-up, the subject had recovered fully, with the exception of the lost vision.

Written informed consent was obtained from the patient for publication of the present case report and any accompanying images.

Discussion

Sphenoid disease is primarily a result of the extension of disease anteriorly from the ethmoid complex or posteriorly from the adjacent anatomical sites, including the sella turcica and the surrounding skull base structures (8). Isolated sphenoid sinus disease is rarely encountered and the associated symptoms are generally non-specific (8). Sphenoid disease frequently leads to nerve involvement, and the effects of masses causing compression, destruction or invasion lead to varying levels of impact on the extent and degree of neurological deficits (8). Etiologically, non-neoplastic lesions constitute the majority of cases, while fungal processes are the third most common cause (8). ISSA is a type of fungal sinus infection that is distinct from other categories of fungal rhinosinusitis, and is characterized by higher levels of occurrence in older women (9). A sphenoid sinusotomy is regarded to be the primary therapy (8,9). To the best of our knowledge, the occurrence of an OH of the...
sphenoid sinus following ISSA surgery has not been previously reported (2-6,8,9).

SNOH is rarely encountered in clinical practice (2-6). When it does occur, it is usually locally invasive and demonstrates a high tendency to occur in the maxillary sinus (3,5,6,10). The incidence of SNOH appears to be most frequent in the population of East Asia (3,5,6,10). The exact underlying etiological mechanisms remain to be elucidated (4,5,6). Ozaki et al (11) advanced what is now the most widely-accepted hypothesis, the negative spiral theory (3,6). The negative spiral theory assumes that an OH results from persistent negative intraluminal pressure following an initial episodic hemorrhage into a semi-closed cavity, leading to repeated rupturing of fragile mucosal vessels. Subsequent formation of a superficial fibrotic capsule prevents further reabsorption of the hematoma. Under the succession of biological healing processes, an OH emerges (3,6). The progressive expansion of the OH results in augmentation of the sinus antrum, and therefore results in demineralization of the adjacent skeletal structures (3,5).

Macroscopically, SNOH is described as a slow-growing, well-circumscribed and friable brownish mass. The histopathological findings reveal a mixture of neovascularization, fibrosis, hemorrhaging and hemosiderin deposition (5). A number of studies have attempted to identify an association between SNOH and sinonasal angiomatous polyps, a rare subtype of inflammatory sinonasal polyp, using radiology and histopathology (10,12). However, this association remains to be fully elucidated. Returning to the present case, the causative etiology may be attributed to an obstructed sinus cavity with equivalent volume to the maxillary sinus, and repetitive post-operative hemorrhaging due to an inflammatory vascular injury. To the best of our knowledge, such a case was previously unknown, and has not been reported in the relevant literature (5). It may be postulated that the incidence of post-operative SNOH is underestimated. Long-term follow-up for patients undergoing endoscopic sinus surgery is necessary to provide an answer to this hypothesis.

CT possesses an advantage over MRI in terms of its ability to analyze the integrity of bone in detail (5). The density in unenhanced CT scans of OH is frequently hyperattenuated.
compared with masticator muscles. The hallmarks of OH are mucoperiosteal thickening, occasional calcification, convex bowing of natural skeletal architecture, cortical thinning or direct extension to adjacent structures sparing frank osseous destruction (5, 6). However, CT alone does not provide enough information to allow for the differentiation of OH from locally aggressive neoplasms. MRI exhibits marked superiority over CT in terms of its ability to determine the margin and true extent of tumor expansion. In addition, MRI possesses an advantage with regard to distinguishing adjacent secondary inflammation and nasal secretions from a tumor mass, and has the ability to display the corresponding pathological components within the lesion (2, 5). Typically, SNOH exhibits a mosaic of varying signal intensities in T1WI and T2WI, and heterogeneous enhancement in a patchy pattern following contrast administration (2, 5, 10). For SNOH, the most conclusive diagnostic finding is a hypointense zone surrounding the lesion on T2WI, indicating the pathological feature of a fibrous capsule (2, 5, 10). Therefore, obtaining knowledge of the distinctive characteristic observations acquired using CT and MRI (2, 5, 10) offers clinicians valuable anatomical and diagnostic informative clues that may allow for the achievement of a correct diagnosis of SNOH.

To the best of our knowledge, the present study reports the first case of an OH of the sphenoid sinus in a patient who previously underwent ISSA surgery. The case highlights the requirement for physicians to include SNOH in the differential diagnosis for expansile sphenoid sinus disease, regardless of its low prevalence. Advances in techniques mean that the performance of endoscopic endonasal skull base surgery is clinically feasible and reliable for the treatment of patients exhibiting SNOH (7), particularly for cases in which patients exhibit compressive neuropathy, and timely intervention is required in order to avoid permanent sequelae (13). The present case additionally suggests that a preexisting inflammatory process and performance of previous surgery may have a role in multifaceted SNOH pathogenesis. However, increased experience with regard to the diagnosis and treatment of sphenoid SNOH is required in order to fully elucidate the underlying mechanisms of the disease.

In conclusion, although the pathogenesis and epidemiology of SNOH are not fully understood, the application of CT and MRI, combined with the involvement of a diagnosing physician who possesses familiarity with the classical imaging findings, allows for an accurate pre-operative diagnosis of SNOH. Care should be exercised when interpreting the abnormal imaging results acquired from a sinus that has been operated on, in order to ensure that SNOH is not misidentified as recurrence or residual disease. Once a diagnosis has been reached, following consultation of the CT and MRI scans, the current understanding is that a complete surgical excision is the optimal treatment for SNOH, yielding positive patient outcomes and rarely resulting in recurrence later in life.

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