Variations in Lateral Sphenoid Sinus Wall Defects

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Abstract

Objective The article aims to present different forms of lateral sphenoid sinus wall defects.

Study Design Case series and literature review.

Methods A comparison between two patients who presented with spontaneous CSF rhinorrhea, defects in the lateral wall of the sphenoid sinus, and meningeal contents in the sphenoid sinus based on MRI is discussed. Both patients were operated endoscopically, through a trans-nasal-ethmoid-pterygoid approach.

Results In the first patient, a meningoencephalocele, protruding through a defect in the lateral sphenoid sinus wall and pterygoid base, occupied the inferior part of the sphenoid sinus. In the second patient, there were no exposed meningeal contents inside the sphenoid sinus. Instead, the lateral sphenoid sinus wall was thin and bulging medially into the inferior part of the sinus, with the dura and temporal lobe covered with thin bony cap.

Conclusion Complete dehiscence of the lateral wall of the sphenoid sinus results in exposed meningeal contents inside the sphenoid sinus. In partial dehiscence, the thin and weakened lateral wall of the sphenoid sinus yields to the pressure of the temporal lobe and bulges into the sphenoid sinus cavity. There are no other descriptions of partially dehiscent lateral sphenoid sinus walls in the literature.

Keywords ► sphenoid sinus ► lateral wall defect ► meningoencephalocele

Introduction

Defects in the lateral wall of the sphenoid sinus are a rare cause of cerebrospinal fluid (CSF) rhinorrhea. The etiology of these defects may be congenital or acquired. A dehiscent lateral cranioharyngeal canal, or Sternberg canal, arises from a malunion of two parts of the developing sphenoid bone: the greater wing and the basisphenoid.1,2 Originally reported in 4% of healthy adults,3 its true prevalence is not known, as is its relation with adult-onset CSF rhinorrhea and meningoceles of the sphenoid sinus. Acquired conditions probably account for the larger part of lateral sphenoid sinus wall defects, and include trauma, adjacent tumor, idiopathic intracranial hypertension, and iatrogenic damage.

Regardless of their cause, large defects in the lateral sphenoid sinus wall are often accompanied by a temporal lobe meningocele or meningoencephalocele. When the defect is small, only CSF would leak, without dural herniation.4 In both cases, the defect should be repaired, achieving high rates of success with both transcranial and endoscopic approaches.5–9

Ongoing pressure plays a role in the pathogenesis of lateral sphenoid sinus wall defects. We hypothesize that apart from complete dehiscence, cases of weakened partially dehiscent walls, bulging into the sphenoid sinus cavity, with micro-tears allowing for CSF leak but without exposed meningeal contents, should also exist. This entity has not been described in the literature so far.
Methods

A comparison of two patients presented with idiopathic CSF rhinorrhea confirmed by positive β₂-transferrin tests is discussed. In both patients, computed tomography (CT) showed an apparent defect of the lateral sphenoid sinus wall, with no underlying extensive pneumatization of the lateral recess. The magnetic resonance imaging (MRI) demonstrated intracranial contents in the sphenoid sinus cavity. Both patients underwent endoscopic surgery, with a trans-nasal-ethmoid-pterygoid approach to the lateral recess of the sphenoid sinus, and multilayer reconstruction of the bony defect. Both patients had continuous lumbar drainage intra- and postoperatively.

Results

Patient 1
A 57-year-old woman, overweight but otherwise healthy, presented with a 6-month history of spontaneous clear rhinorrhea, with no previous head trauma or surgery. During that period she suffered from two episodes of pneumococcal meningitis, which were successfully treated with intravenous (IV) ceftriaxone each time. CSF leakage was confirmed by a positive β₂-transferrin test. A high-resolution CT demonstrated total opacification of both sphenoid sinuses, a 6-mm defect in the lateral wall of the left sphenoid sinus, and a 9-mm defect in the roof of the pterygoid base (► Fig. 1). T2-weighted MRI images showed a temporal lobe meningoencephalocele occupying the lower half of the left sphenoid sinus and the base of the pterygoid plates (► Fig. 2). An endoscopic trans-nasal-ethmoid-pterygoid approach was chosen to open the entire left sphenoid sinus. After removal of the sphenoidal rostrum and dissection of the pterygopalatine fossa, the meningoencephalocele was exposed (► Fig. 3). The latter was shrunk by electrocautery and resected at its base, and the bony defect was sealed by a multilayer reconstruction using an abdominal fat plug and an overlying free septal mucosal graft.

Patient 2
A 43-year-old otherwise healthy man presented with an uncomplicated 3-week-old episode of spontaneous left rhinorrhea approved as CSF leak by means of β₂-transferrin test. There was no previous history of head trauma or surgery. His high-resolution CT scan revealed a 10-mm defect in the inferior part of the lateral wall of the left sphenoid sinus and a bony septum traversing the sinus cavity. The left sphenoid sinus was totally opacified, whereas the right sphenoid sinus was partially obstructed. The
pterygoid plates were intact bilaterally (► Fig. 4). MRI demonstrated meningeal contents in the left sphenoid sinus (► Fig. 5). Similar to the first patient, a continuous lumbar drainage was installed, the left sphenoid sinus was reached by an endoscopic trans-nasal-ethmoid-pterygoid approach. Unlike the first patient, after the entire left sphenoid sinus cavity was revealed, we found no exposed intracranial contents, but rather a bony bulge occupying the inferolateral part of the sinus. No frank CSF leak was evident. Carefully breaching that thin layer of bone, we revealed the middle fossa dura underneath (► Fig. 6). The entire bony shell was removed, and the meningocele shrunk by electrocautery and resected at its base. The 1-cm large lateral wall defect was then bridged in a multilayer fashion, using septal cartilage and a contralateral nasoseptal vascularized flap.

Discussion

The pathogenesis of defects in the lateral wall of the sphenoid sinus is complex. In some cases, such as trauma or surgery, the bony wall is disrupted by an abrupt insult. In other cases involving elevated intracranial pressure, progressive pressure gradually weakens the bone, ultimately creating a complete defect. In pure congenital cases, a developmental error in the ossification of different parts of the sphenoid bone results in a malformed lateral wall of the sphenoid sinus. Pathogenesis can be multifactorial, as proposed by Tabae et al, when an asymptomatic congenital base of skull defect enlarges over time by continuous CSF pulsations, until the development of a dural tear, a meningoencephalocele, or both.5

Both transcranial and endoscopic approaches were described to seal the CSF leak and reconstruct the skull base defect. Evidence supports applicability of an endoscopic approach for most cases, reserving transcranial surgery for recurrent cases after failed endoscopic attempts.2,5–8

The rising number of reports in recent years suggests that lateral sphenoid sinus wall defects are perhaps not as rare as previously thought.5–7,9 In these reports, the sphenoid sinus is partially or entirely occupied by a meningoencephalocele, and the lateral wall of the sphenoid sinus is absent.

Similar findings could be seen in our patient 1. ► Fig. 3 shows the surgically exposed left sphenoid sinus occupied in its inferior part by a meningoencephalocele, originating from a fully dehiscent lateral sphenoid sinus wall.

Vaezi et al have described a different version of lateral sphenoid sinus wall defects, in which CSF collections or meningoceles were covered by normal sphenoid sinus mucosa.4 The lateral wall of the sphenoid sinus in these cases was either absent, accompanied by a meningocele, or intact with assumed micro-tears, with CSF collections.

Although two of the main pathogenetic pathways of lateral sphenoid sinus wall anomalies, congenital maldevelopment
and gradual pressure, would result in a weak and thin lateral wall, descriptions of thin lateral walls bulging into the sphenoid sinus cavity are absent in the literature.

In our patient 2, preoperative imaging supported the diagnosis of a lateral sphenoid sinus wall dehiscence with an accompanying meningoencephalocele in the sphenoid sinus. However, intraoperatively, we found a thin medially displaced lateral sphenoid sinus wall, without dural exposure (► Fig. 6). Because intracranial pressure (ICP) was lowered intraoperatively by a continuous lumbar drain, there was no evident CSF leak in the operative field. We believe that although there was no apparent dehiscence of the skull base, CSF leaked to the nose through microscopic defects in the thin medially bulging lateral wall. Removing this bony cap and the dural contents behind it confirmed the absence of a normal lateral sphenoid sinus wall. When we reviewed the patient’s preoperative CT scan, the medially displaced lateral sphenoid sinus wall was visible as the transverse septa crossing the sinus cavity (► Fig. 4).

**Conclusion**

When evaluating a patient with an apparent meningoencephalocele in the sphenoid sinus, surgeons should take into consideration the possibility of encountering a partially, rather than completely, dehiscent lateral sphenoid sinus wall.

**References**