MRI and CT Imaging of an Intrasphenoidal Encephalocele: A Case Report

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Summary

Background: Intrasphenoidal encephalocele (ISE) is a rare clinical entity. The incidence of congenital encephalocele is very low. Accurate diagnosis and surgical approach is of critical value.

Case Reports: We present a case of intrasphenoidal encephalocele in a 40-year-old man. He complained of cerebrospinal fluid (CSF) rhinorrhea and recurrent meningitis. In images of computed tomography (CT) and magnetic resonance imaging (MRI), intrasphenoidal encephalocele herniating through a defect of the left lateral sphenoid sinus wall was determined. Incisional biopsies were taken by endoscopic transnasal approach and histopathological examination revealed an encephalocele. In the differential diagnosis, ISE can be taken for inflammatory or malignant sinusoidal soft tissue masses. ISE is differentiated from other entities by demonstrating continuity with normal brain tissue.

Conclusions: MRI clearly demonstrates that the herniating soft tissue is isointense with brain and continuous with brain tissue via the sphenoid sinus, thereby the treatment decision-making process is very important.

MeSH Keywords: Encephalocele • Magnetic Resonance Imaging • Multidetector Computed Tomography

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the defect was seen in the left sphenoid sinus (Figure 1). What is more, bilateral, apparent petrous apex pneumatization was noticed. In order to identify the structure of the polypoid mass in the left sphenoid sinus, brain magnetic resonance imaging (MRI) with contrasting agent was obtained. In brain MRI, intrasphenoidal encephalocele was seen (Figure 2). With those findings, the patient underwent transnasal endoscopic surgery. Incisional biopsy was taken from the polypoid mass in the sphenoid sinus (Figure 3).

The histological examination of the specimens taken from the sphenoid sinus revealed glial tissue in fibrovascular connective tissue fragments of mucosa coated by respiratory-type epithelium (Figure 4A). Immunohistochemical stains for glial fibrillary acidic protein (GFAP) and S-100 confirmed the diagnosis (Figure 4B).

**Discussion**

Intrasphenoidal encephalocele (ISE) is a rare clinical condition. The incidence of congenital encephalocele is approximately 1 in 3000–5000 live birth [5]. ISE has an estimated incidence of 1 in 700,000 live birth [6]. A small occult congenital dysplasia in the form of defects or clefts in the base of the skull in the middle cranial fossa is suggested in the etiology of ISE. These structures constituting a weak area, under the pressure of the CSF may favor the creation of meningoceles or encephaloceles which finally become clinically apparent as spontaneous CSF rhinorrhea [7]. Incomplete posterior portion of the bony fusion in the sphenoid sinus creates a lateral craniofaryngeal canal (Sternberg’s canal). Sternberg’s canal is attributed
as a possible site of origin of congenital encephalocele. Sphenoidal defects at fusion planes are more likely to be congenital than acquired [3].

Intracranial hypertension, by means of arachnoid pits in the lateral recess of the sphenoid sinus, is the major cause of acquired spontaneous CSF leaks and lateral intrasphenoidal encephaloceles [8]. Intrasphenoidal encephaloceles that have an intact dura and no CSF leak are generally reported during imaging studies for other causes [9].

The CSF volume is generally not affected because of intermittent drainage. Recurrent meningitis may also occur due to CSF drainage. Although there were both rhinorrhea and recurrent meningitis in the presented case, there was no history of trauma or tumour. We suggested that the bone defect in the lateral wall of the sphenoid sinus could be Sternberg’s canal.

In radiological imaging of encephalocele, CT, a non-invasive technique, shows bone structures in details and is more precise in detecting the defect sites in the skull base. Bone defect in the lateral wall of the left sphenoid sinus and soft tissue beside the defect and with the same density as the brain tissue were noticed in CT images of the presented case. In the differential diagnosis of soft tissue of that kind, inflammatory and tumoral lesions originating from the sphenoid sinus and cerebral tissue should be considered initially. Due to the fact that MRI is more accurate in visualizing soft tissues in detail, it should be performed. In the presented case, continuity of the cerebral tissue in the left sphenoid sinus was noticed in MRI and it was regarded as ISE (Figure 2A, 2B). Biopsy was carried out by transnasal endoscopic approach and soft tissue sample was subjected to histopathological examination. The differential diagnosis of ISE includes nasal teratoma, glial heterotopia and also a true glioma. ISE should be distinguished from other developmental anomalies and cystic teratomas based on characteristic clinical and radiological findings [10].

Because a persistent CSF leak may lead to meningitis or brain abscess, it is potentially lethal. Therefore, intrasphenoidal encephalocele should be repaired by surgical intervention. We used transnasal endoscopic approach in the presented case. After pushing the brain tissue toward the endocranium, temporal fascia and a free muscle graft were inserted into the defect of the lateral wall of the left sphenoid sinus and then fixed with fibrin glue.

Conclusions

Intrasphenoidal encephalocele often associated with a persistent Sternberg’s canal is a rare entity. Our patient presented with recurrent meningitis and CSF rhinorrhea. Malignancy, inflammatory changes and encephalocele are taken into account in the differential diagnosis of bone defect seen in CT. MRI examination shows cerebral tissue that extends into the sphenoid sinus. MR examination is extremely important in diagnosing intrasphenoidal encephalocele. Knowing that the soft tissue within the sphenoid sinus is encephalocele, allows us to change the therapeutic approach.

Conflicts of interest

The authors declared no conflicts of interest.

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