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Authors: Özgür Ertuğrul, Esra Çiçekçi, Mehmet Cudi Tuncer, Mehmet Ufuk Aluçlu

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Recurrence of atretic parietal cephalocele in adult: report a case and review of literature

Short title: Atretic parietal cephalocele

Özgür Ertuğrul¹, Esra Çiçekçi², Mehmet Cudi Tuncer³, Mehmet Ufuk Aluçlu⁴
¹Department of Radiology, Private Bağlar Hospital, Diyarbakır, Turkey
²Department of Physiotherapy, University of Health Sciences, Gazi Yaşargil Education and Research Hospital, Diyarbakır, Turkey
³Department of Anatomy, Faculty of Medicine, University of Dicle, Diyarbakır, Turkey
⁴Department of Neurology, Faculty of Medicine, University of Dicle, Diyarbakır, Turkey

Address for correspondence: Mehmet Cudi Tuncer, Professor, PhD, Department of Anatomy, Faculty of Medicine, University of Dicle, Diyarbakır, Turkey, tel: +90 412 2488001 Ext. 4539 (Faculty room), fax: +90 532 2744926 (Mobile phone), e-mail: drcudi@hotmail.com

ABSTRACT

Common presentation of atretic parietal cephalocele (APC) is mostly seen in infants and young children. And, it is a palpable midline parietal soft tissue mass which is thought to represent involuted true cephalocele (meningocele or encephalocele) connected to dura mater via a fibrous stalk. The incidence of intracranial anomalies have increased with APC. Parietal cephaloceles comprise about 1% of all cerebrospinal congenital malformations and 10% of cephaloceles. We report here the case of an atretic parietal cephalocele with no associated brain malformations in adult.

Key words: atretic cephalocele, encephalocoele, straight sinus, brain herniation, adult, neuroradiology

INTRODUCTION

Atretic parietal cephalocele, also known as atretic cephalocele (AC), refers to small subscalp lesions that consist of dura, fibrous tissue, neural and glial rests [41]. Cephaloceles
typically present at birth and usually come to medical attention within the first week of life. They are categorized according to the site of the skull defect. They are often sporadic, but the occipital form may be part of a genetic malformation syndrome. Most cephaloceles and associated malformations are treated by surgery. The prognosis depends on the site, content of the sac, operability, hydrocephalus, and other associated malformations. Cephaloceles are designated as: occipito-cervical type, if the skull defect involves the occipital bone, foramen magnum and the posterior arches of C1, C2, etc.; occipital type, if the defect lies between foramen magnum and the lambda; parietal type, if the defect lies between the lambda and the bregma; lateral type, if the defect lies along the coronal or lambdoid sutures as far inferiorly as the anterolateral and posterolateral fontanelles; interfrontal type, if the defect lies between the bregma and the nasal bones; temporal, if the defect lies along the superior surface of the petrous pyramid; fronto-ethmoidal type (synonym: sincipital), if the defect lies between the nasal bones and the ethmoid bone; sphenoid type, if the ostium for the herniation involves the optic foramen, the superior orbital fissure, or a defect in the orbital wall; sphenomaxillary type, if the ostium for the herniation extends through the superior orbital fissure and the inferior orbital fissure into the pterygopalatine fossa; nasopharyngeal type, if the defect lies within the ethmoid, sphenoid, or basioccipital bones [48]. Martinez-Lage et al. [32,33,34] classified AC into two types: Type 1 consisting of arachnoid tissue with clusters of anomalous blood vessels limited to the stalk of the lesion, Type 2 consisting of meningeal tissue intermingled with dermal and fibrous tissue, as well as clusters of anomalous blood vessels, extending as a net, and ectopic neural or glial elements. In according to classification of Martinez-Lage et al. [32,33,34], our case can be evaluated as type 1. Because, it is clear that localized soft tissue mass under the scalp (Figure1,2 and 3). In addition, Peters et al. detected subscalp lesions that are covered by skin [43]. Atretic parietal cephaloceles differ from the other cephaloceles as defined by Yokota by having small, non-cystic, flat or nodular lesions arising in the vertex midline [54]. Magnetic resonance imaging (MRI) and computed tomography (CT) are the best imaging tools to evaluate the possible concomitant central nervous system (CNS) anomalies in patients with atreticcephaloceles. Pathological classification of cephaloceles is known as meningocele (leptomeninges and cerebro spinal fluid-CSF), meningoencephalocele (leptomeninges, CSF and brain), meningoencephalocele (leptomeninges, CSF, brain and ventricles), atretic cephalocele (small nodule of fibrous fatty tissue), gliocele (CSF lined by glial tissue) [7,39]. Atretic parietal cephaloceles are associated with anomalies of Chiari II and III malformations, corpus callosum agenesis and Dandy-Walker malformation [32,38].
The etiology of cephalocele in the majority of cases is not known. Like neural tube defects, cephalocele is considered to be multifactorial in origin, and a variety of genetic and environmental factors have been implicated in its etiology [12,22]. The influence of genetic factors is suggested in some cases of occipital cephaloceles that are associated with genetic syndromes such as Knobloch syndrome [46], Meckel syndrome [40], and other rare syndromes [14]. Moreover, the significance of the role of genetic factors in occipital cephalocele may be supported by its increased incidence in families with previous neural tube defects [56]. The incidence of cephaloceles is approximately 0.8-3.0 per 10,000 births [20,28,48]. Thus, cephaloceles are approximately 10 times less common than myelomeningoceles. Mealey et al. [36] found 60 encephaloceles and 559 myelomeningoceles among 76,280 admissions to a children's hospital over a 20 year period. The different types of cephaloceles show strong geographic variation, substantial differences in incidence and form with race and sex, as well as different associations with neural tube defects: among white populations in Europe and North America, occipital cephaloceles constitute 66%-89% of all cephaloceles [13,48]. In Southeast Asia, sincipital cephaloceles are more common [50]. Among white Australians, 66.7% of cephaloceles are occipital and only 2.2% sincipital, whereas among aboriginal Australians, 50% of cephaloceles are sincipital [48]. Igbos Nigerians have a similar, disproportionately high incidence of sincipital cephaloceles. It is clearly seen that genetic origin is important to define different types of cephaloceles. The male to female ratios are 1/(2.0-2.7), approximately 2:1, and approximately 1.5/1, respectively. Occipital and parietal cephaloceles appear to be related to neural tube defects [48]. Mealey et al. reported that 7% of children with occipital encephalocele show concurrent myelomeningocele [36]. Three percent of patients with (cervico)occipital cephalocele have diastematomyelia and 3% have a second cephalocele in the parietal region the same as our in patient [48].

The diagnosis is based on radiological findings. Ultrasonography is of great value in determining the size and content of the cephalocele, the size and shape of the skull, the extent of bony defect, and the size and shape of the ventricular system and its relation to the cephalocele sac in both the fetus and the neonate [11,27]. This method is particularly useful in the prenatal diagnosis of cephalocele and associated malformations [8,24,44,45,47]. In the postnatal period, the CT scan and MRI provide us with the most useful information regarding anatomic structure in delineating various brain and skull abnormalities associated with the occipital cephalocele [55,57].
We report here a case of a recurrent atretic parietal cephalocele, which is rarely seen, with characteristic imaging features in adult. We described all types of AC in detail and discussed pathological classifications of cephalocele in this review.

CASE PRESENTATION

A 25-year-old man who had been operated for a subscalp mass when he was 5 years old. Twenty years later, he was admitted again to neurosurgery clinic for recurrence of the mass on the operation site at a rural hospital. The pathology of mass was normal in his report, and excision of superficial mass was misdiagnosed as dermoid cyst. Over two years, the patient had been complaining of gradually increasing mass on the operation site. Therefore, the patient was admitted to neurosurgery clinic in our university. In routine radiological examinations, CT and MRI of the brain were performed for characterization of the mass. A small cranial defect at the posterior midline of parietal bone which was connected to the scalp mass was observed on noncontrast CT (Figure 1A and 1B). The lesion was hyperintense on T2 W and hypointense on T1 W images. Sagittal T2W images demonstrated a bony defect involving the underlying parietal bone. T2W sagittal and axial images showed flow void extending posterosuperiorly representing embryonic position of the vertical straight sinus and prominent superior cerebellar recess (Figure 2).

Based on CT and MR imaging findings, we present here recurrent atretic parietal cephalocele in adult which is not associated with malformations of APC (Chiari II and III malformation, corpus callosum agenesis and Dandy-Walker malformation etc.).

DISCUSSION

Atretic cephaloceles are formes frustes of cephaloceles that are characterized by a small, noncystic, flat, or nodular lesion situated in the midline of the scalp, either near to vertex (parietal form) or just cephalic to the external occipital protuberance (occipital form) [54]. The prognosis for patients with a cephalocele depends primarily on site, content of the sac, operability, hydrocephalus, and additional malformations [16,25,29,30,31,34,36,48,53].

Historically, the term cephalocele, in a restrictive sense, is defined as a protrusion of part of the cranial contents through a congenital opening in the cranium, typically covered
with skin or mucous membrane. However, in a broad sense of the term, it has been used to include also the acquired traumatic and nontraumatic forms. The condition was first described by Forestus in 1590 [6]. The first monograph on the subject was written by Corvenius in 1749, and was followed by a long series of papers on the various aspects of the disorder including the clinical features, pathology, incidence, pathogenesis, and treatment [6].

Ballantyne's comprehensive morphologic study contributed significantly to the understanding of the pathology of the lesion and its classification. A variety of terms have been used to describe this congenital lesion. They include "hernia cerebri," "hydranencephalocele," "encephalocele," "cephalocele," "cranium bifidum," "exencephalus," and "exencephalocele." The term "hernia cerebri" should be reserved for acquired lesions. "Hydranencephalocele" is confused with "hydranencephaly," which represents a different congenital lesion [21]. "Cranium bifidum" describes only the bony defect. "Exencephalus" is confused with "exencephaly," which represents an earlier stage of anencephaly [56]. The term "exencephalocele" is also confusing and should be avoided. Among these, the term "cephalocele" appears to be the most appropriate [6,17]. This term denotes 2 specific types of lesions: herniation of brain tissue with the overlying meninges through a cranial defect (meningoencephalocele or encephalocele); and herniation of meninges without brain tissue (cranial meningocele). Today, the terms cephalocele and encephalocele are used almost interchangeably in the neurosurgical literature. Diminutive forms of cephalocele appear as small noncystic subcutaneous nodules and are often of no clinical significance. These forms have been referred to under a variety of terms including "occult," "rudimentary," "abortive," "sequestrated," and "atretic" encephalocele or meningocele of scalp [34,41,54]. The other terms used for these subcutaneous lesions include "brain tissue heterotopia," "glial heterotopia," and "glioma" in scalp and nasal regions [15,51].

Atretic encephalocele was first reported in 1972 by James and Lassman, who defined it as a degenerative form of encephalocele [26]. Its reported incidence is 4-17% of all the encephaleles [54] and parietal location comprises 40-50% of cases. It has been rarely reported in adults [2]. Parietal cephaloceles are very rarely seen (1% of all cerebrospinal malformations and 10% of cephaloceles [43] and if they are congenital, they are usually associated with many anomalies such as corpus calosum agenesis, Chiari II, Dandy-Walker malformation. McLaurin [35], in his series of parietal cephalocele, noted that 3 of 4 patients with encephalocele had major developmental brain abnormalities and died; the fourth patient remained alive but with severe psychomotor retardation associated with phenylketonuria. Of 7 cases of parietal meningocele, only 2 had brain abnormality that consisted of agenesis of the
corpus callosum, and no one of them died during 9 months to 10 years of follow-up. Simpson et al. noted that among 8 cases of parietal encephalocele that were followed up, 7 were severely handicapped, but none of the 3 parietal meningocele patients showed significant disability [48]. In a study by Patterson et al., 5 of 8 patients had parietal AC as an isolated anomaly, while 3 patients had other brain abnormalities, with the most common additional finding being ventriculomegaly [41].

Generally, venous anomalies accompany this disease, and most of these cases have benign clinical findings. Causation of abnormal venous drainage in the great vein of Galen, the straight sinus and the superior sagittal sinus may be secondary, through interaction with a developmentally pre-existing encephalocele [23,37]. Regardless of the mechanism of atretic cephalocele formation, vertical embryonic positioning of the straight sinus has frequently been identified in these lesions and deserve to be mentioned both as a marker of the timing of the embryologic insult and as a clue to radiologic diagnosis "hallmark imaging sign". During stage 7a of cranial venous development, corresponding to a crown-rump length of approximately 80 mm, the straight sinus is nearly vertical in a ventral-dorsal plane. We also determined that the straight sinus was coursing vertically and prominent superior cerebellar recess in T2W sagittal and axial images (Figure 2). Various malformations may be seen in APC, especially in Chiari II and III malformations, corpus callosum agenesis and Dandy-Walker malformations [19,37]. Other associated malformations include ventriculomegaly, cystic malformations, heterotopias of the posterior fossa. In this case, we have not seen any concomitant anomaly except for the parietal location and typical imaging findings.

The bone deformities of AC could be seen on plain x-rays [5,10]. However, not all patients with AC present with bony defects or scalp lesions [18]. MRI and CT are the best imaging modalities to evaluate the possible concomitant CNS anomalies in patients with AC. The bony margins of an AC could be oval or elongated in shape, and taper from inside to outside. The cystic content is isodense to the cerebrospinal fluid or slightly hyper-dense to the brain parenchyma in CT [1,5,32]. The cyst is hypointense in T1- and hyperintense in T2-weighted MRI scans [3,9]. The AC content enhances if a vascular structure is present inside it [1]. We observed that the lesion was hyperintense on T2 W and hypointense on T1 W images as the same results in other studies [3,9].

CONCLUSIONS

In conclusion, we present here recurrent APC, which is not associated with brain malformations in adult. The incidence of APC is rarely seen in adults. A few case reports of
APC in adults have been reported in literatures so far [1,2,4,5,23,42,49]. The most important diagnostic workups include ultrasonography for prenatal diagnosis and MRI and CT in children and adults [27,52]. The presence of straight sinus coursing vertically and prominent superior cerebellar recess is detected through radiological findings. Especially, the straight sinus coursing vertically is the most important landmark to radiologically define APC.

REFERENCES


FIGURES

**Figure 1A.** A midline small subscalp mass on unenhanced CT (red asterisk)

**Figure 1B.** A small bone defect (blue asterisk) at the posterior midline of parietal bone which was connected to the scalp mass (red asterisk)
Figure 2. Sagittal T2W images demonstrated flow void extending posterosuperiorly representing embryonic position of the vertical straight sinus (long black arrows), prominent superior cerebellar recess (red asterix) and a small subscalp mass (black arrow).