Coexistence of Temporo-Occipital Polymicrogyria with Choroidal Fissure Cyst in a Case of Focal Onset Seizures

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Abstract

Focal-onset seizures can be caused by underlying brain lesions including focal lesions such as granulomas, low-grade neoplasms, vascular lesions, or neuronal migration disorders. Polymicrogyria is a congenital abnormality of cortical formation occurring during embryonic life. Choroidal fissure cysts are either arachnoid or neuroepithelial cysts arising at the choroidal fissure, and mostly they are incidental findings having no significant clinical implications. Coexistence of both of these can lead to dilemma in the management decisions. We present a case of focal-onset seizures with an unreported coexistence of polymicrogyria with choroidal fissure cyst.

Keywords

► polymicrogyria
► choroidal fissure cyst
► seizures
► MRI

Introduction

Focal lesions such as granulomas, low-grade neoplasms, vascular lesions, or neuronal migration disorders can be causative factors in adult-onset seizures.¹ Polymicrogyria (PMG) is an abnormality of fissures and sulcation due to various insults occurring during cortical development in embryonic life.² It may manifest as seizures and usually diagnosed early in life with the help of magnetic resonance imaging (MRI). Choroid fissure cysts (CFCs) are mostly incidental findings in brain MRI of general population, but may be of significance in those presenting with seizures as previous reports have mentioned their association with complex partial seizures.³⁻⁵

Case Report

A 20-year-old male patient presented with recurrent headaches and seizures since 6 years of age. He was prescribed an antiepileptic drug by a practitioner near his home but continued to have seizures. His seizures were preceded by an aura of vertigo with visual disturbances in both eyes followed by eye deviation, neck turning to left side, oral automatisms, impaired awareness of the episode, and bimanual posturing. But there was no loss of posture or bilateral spread during event. There was history of nocturnal events and sudden on-and-off events. No postictal phenomenon or memory was present for the event. There was history of unresponsiveness and behavioral arrest during the event. The last episode was a day prior to presentation to the hospital. His intellectual and behavioral features were normal. Systemic examination was normal. Electroencephalogram revealed abnormal interictal epileptiform discharges in right posterior leads, with generalized slow wave discharges. He was diagnosed to have focal nonmotor onset sensory (visual) seizures with impaired awareness and was referred for imaging. Plain computed tomography brain revealed fluid-density lesion lateral to midbrain in the right choroid fissure region. MRI of the brain (plain and contrast) was performed. A well-defined lobulated cerebrospinal fluid (CSF) intensity lesion (~Fig. 1) was seen medial to mediotemporal lobe on right side. There was a mass effect in the form of compression with associated partial inversion of hippocampus. Mild prominence of the temporal horn of ipsilateral lateral ventricle was noted with lateral displacement of choroid plexus. There was no diffusion restriction or contrast enhancement suggesting CFC. Multiple small gyri and sulci are noted in ipsilateral temporo-occipital lobe with normal signal as compared with rest of the brain parenchyma suggesting PMG (~Fig. 2).
Discussion

Routine neuroimaging is recommended for all patients having seizures, particularly those in adult-onset and with focal type of seizures, at least once, usually before starting antiepileptic therapy. The likelihood of underlying lesions is more in those with refractory seizures and may benefit from MR neuroimaging to delineate lesion for management and if necessary, by surgical planning. PMG is a neuronal developmental disorder characterized by presence of multiple small partly fused gyri and sulci, and irregularly appearing cortical surface as suggested by its name. It is due to abnormal cortical formation resulting from disturbance in cortical development late in the neuronal migration stage or early in the cortical organization stage. Causes are multifactorial ranging from prenatal infection, ischemia, or exposure to toxins, to chromosomal abnormalities. The common patterns of PMG are perisylvian (61%) with parasagittal parieto-occipital accounting for 3%, and there is association with periventricular gray matter heterotopias in 11%. In perisylvian PMG,
85% are bilateral and are symmetrical. The median age at presentation noted in the former study is 4 months. Nearly more than one-third (38%) of cases were diagnosed in either antenatal or neonatal period. Seizures are the most common clinical sequelae of PMG with approximately 80% of patients eventually developing seizures and majority within the first 5 years. Other symptoms can be encountered depending on the area of the brain involved. Bilateral abnormalities may show various syndromic associations. On MRI, the diagnostic criteria for PMG include unusually thickened and over-folded gray matter, cortical surface irregularity, and “stippling” or irregularity at gray–white matter interface. It appears as multiple cortical convolutions and shallow sulci with thickened or normal cortex. An anomalous vein may occasionally be seen in the region of PMG. MRI is the imaging technique of choice for diagnosing PMG. The management of PMG constitutes antiepileptic drugs and timely follow-up.

Choroid fissure cysts are CSF-like fluid-containing benign cysts that may be congenital in origin. The CFC can be of arachnoid or neuroepithelial origin, differentiated only at pathology. However, most are not confirmed with histopathology as they do not require surgery. There are reports of association of CFC with seizures, attention-deficit hyperactivity disorder, migraine, and narcolepsy. There are only at pathology. However, most are not confirmed with histopathology as they do not require surgery. There are reports of association of CFC with seizures, attention-deficit hyperactivity disorder, migraine, and narcolepsy. They do not show contrast enhancement or adjacent edema or gliosis. CFCs are found incidentally, and the treatment is conservative with interval follow-up, if necessary. If the lesion is large and symptomatic, then surgical treatment includes cyst fenestration or cystoperitoneal shunting.

Usually, the patients of cortical malformations clinically manifest at a much younger age. Our case report emphasizes the importance of looking for more subtle abnormalities such as PMG even in the presence of an obvious lesion of CFC while evaluating an adult patient presenting with seizure for the first time without any history of trauma or headache. Subtle changes of PMG can be missed when associated with a cystic lesion as the cyst itself can have cortex-deforming mass effect as noted in our case. There have been no reported cases showing coexistence of CFC and PMG. The importance of this coexistence lies in treatment decisions as both conditions are associated with seizures. Seizures if associated with only CFCs may resolve with decompression or fenestration of the cyst or shunting. The presence of associated PMG may warrant long-term treatment with antiepileptic drugs if surgical resection is not feasible. In our patient, the patient’s seizures were under control with an antiepileptic drug. This case highlights the dilemma as and when if seizures become refractory in such a patient with dual pathology. The neurosurgery opinion is to perform a decompression or excision of the cyst first, if seizures become refractory, and then to look for control of seizures with medical therapy. The patient is being followed up on an outpatient basis.

Conflict of Interest
None declared.

References