The Perirolandic Sign: A Unique Imaging Finding Observed in Association with Polymerase γ-Related Disorders


AJNR Am J Neuroradiol 2020, 41 (5) 917-922
doi: https://doi.org/10.3174/ajnr.A6514
http://www.ajnr.org/content/41/5/917
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ABSTRACT

SUMMARY: Pathogenic variants in the polymerase γ gene (POLG) cause a diverse group of pathologies known as POLG-related disorders. In this report, we describe brain MR imaging findings and electroencephalogram correlates of 13 children with POLG-related disorders at diagnosis and follow-up. At diagnosis, all patients had seizures and 12 had abnormal MR imaging findings. The most common imaging findings were unilateral or bilateral perirolandic (54%) and unilateral or bilateral thalamic signal changes (77%). Association of epilepsia partialis continua with perirolandic and thalamic signal changes was present in 86% and 70% of the patients, respectively. The occipital lobe was affected in 2 patients. On follow-up, 92% of the patients had disease progression or fatal outcome. Rapid volume loss was seen in 77% of the patients. The occipital lobe (61%) and thalamus (61%) were the most affected brain regions. Perirolandic signal changes and seizures may represent a brain imaging biomarker of early-onset pediatric POLG-related disorders.

ABBREVIATIONS: ASL = arterial spin-labeling; EEG = electroencephalogram; EPC = epilepsia partialis continua; MELAS = mitochondrial encephalomyopathy with lactic acidosis and stroke-like episodes; mtDNA = mitochondrial DNA; Pol γ = polymerase γ; POLG-RD = DNA polymerase γ-related disorder

POLG polymerase γ (Pol γ) is the only DNA polymerase active during human mitochondrial DNA (mtDNA) genome replication. Pathogenic variants in the Pol γ gene (DNA polymerase γ [POLG]) cause a group of clinical syndromes known as POLG-related disorders (POLG-RDs). Patients with POLG-RDs fall into a heterogeneous clinical spectrum. At the least severe end of the spectrum, patients present in adulthood with ptosis and ophthalmoplegia, whereas those most severely affected present with progressive and severe neurologic impairment and liver involvement in early childhood.

Pathogenic variants of POLG are the most frequently detected genetic forms of mitochondrial epilepsy. Seizures are described as the first clinical manifestation in up to 50% of patients. Seizure types include myoclonus, focal motor seizures, generalized seizures, status epilepticus, refractory febrile seizures, and epilepsy partialis continua (EPC).

Neuroimaging findings in POLG-RDs have been described primarily in the later stages of the disease, with the occipital lobe being the most commonly involved region. Brain MR imaging findings in the early onset of pediatric POLG-RDs are not well-known. The primary goal of this study was to describe the early brain MR imaging findings in children, including the “perirolandic sign,” defined as signal changes in the brain parenchyma surrounding the central sulcus of POLG-RDs, and to correlate them with electroencephalogram (EEG) findings. Our secondary goal was to describe the evolution of brain MR imaging findings on follow-up imaging.

Case Series

This retrospective institutional review board–approved study was conducted in a single academic pediatric hospital, Children’s Hospital of Philadelphia. Medical records were searched for primary mitochondrial disorders from January 2001 to July 2018. Patients with brain MR imaging and a confirmed molecular diagnosis of POLG-RDs (confirmed biallelic inheritance for autosomal recessive disease when possible) were included. Exclusion criteria consisted of the unavailability of brain MR imaging or confirmed molecular diagnosis.
findings were pooled and summarized in 1 single column in On-line Table 2, called “Pooled Follow-Up Imaging.”

Most patients were female ($n = 9$, 69%). The median age was 3 years (interquartile range, 0.7–7.5 years). All patients presented clinically with seizures ($n = 13$, 100%). Other common associated symptomatology included regression, developmental delay, hypotonia, vomiting, and signs of liver damage. All patients had an EEG available around the time of their first brain MR imaging ($n = 13$, 100%); all the scans had abnormal findings. Clinical or EEG evidence of EPC was detected in most patients ($n = 8$, 61%). The mean elapsed time between the most contemporaneous EEG and brain MR imaging at diagnosis was 6 days. The most common pathogenic variant found from each parent with another variant was c.1399G>A:p. A467T ($n = 7$, 54%), which is consistent with findings in previous literature.6

At diagnosis, brain MR imaging findings were abnormal in most patients ($n = 12$, 92%). The most common brain MR imaging findings were unilateral or bilateral pericentral signal changes ($n = 7$, 54%) (Fig 1) and unilateral or bilateral thalamic signal changes ($n = 10$, 77%) (Fig 2). Pericentral signal abnormalities were unilateral in most cases ($n = 5$, 71%) and were more frequently seen only affecting the precentral or both the pre- and postcentral gyri. Half of the thalamic changes were unilateral ($n = 5$, 50%). Simultaneous pericentral and thalamic signal changes occurred in 6 patients ($n = 6$, 46%). An association of EPC (clinically/EEG) and pericentral signal changes was present in 6 patients ($n = 6$, 73%), and an association of EPC (clinically/EEG) and thalamic signal changes, in 7 patients ($n = 7$, 87.5%). Two patients with EPC (clinically/EEG) did not present with pericentral signal abnormalities. The findings positive for lesions overall were on the DWI of 10 patients ($n = 10$, 83%), on the T2WI of 9 patients ($n = 9$, 75%), and on FLAIR of 7 patients ($n = 7$, 58%). In 4 patients, DWI was the only sequence with abnormal findings. The occipital lobe was affected in the early brain MR imaging in 2 patients ($n = 2$, 15%). Signal changes in other brain regions were found in 5 patients ($n = 5$, 42%), involving multiple regions, namely the cerebral white matter, insula, putamen, caudate nucleus, fornix, cerebellar vermis, and also the frontal and occipital lobes. One patient did not present with either pericentral or thalamic changes but instead presented with a diffuse pattern similar to that of leukoencephalopathy, with restricted diffusion in the white matter and white matter tracts (Fig 3). Three patients had an abnormally high lactate peak on MR spectroscopy. None of our patients had ASL or other perfusion-weighted imaging at the time.
**DISCUSSION**

Polγ is the only human DNA polymerase active during mtDNA replication.\(^1\) **POLG** is a nuclear gene that encodes for Polγ, and pathogenic variants lead to errors in mtDNA replication, resulting in multiple small mtDNA deletions, an overall reduction in the number of mtDNA genome copies (mtDNA depletion), and decreased adenosine triphosphate production leading to chronic loss of cellular energy.\(^2\)

Patients with **POLG-RD** have been grouped into multiple distinct clinically-defined syndromes, which were later attributed to different subtypes of pathogenic variants. The prototypical clinical phenotypes associated with **POLG** pathogenic variants are the following: 1) Alpers-Huttenlocher syndrome; 2) myocerebrohepatopathy spectrum; 3) myoclonic epilepsy myopathy sensory ataxia; 4) ataxia neuropathy spectrum, which includes mitochondrial recessive ataxia syndrome, spinocerebellar ataxia with epilepsy, and sensory ataxia neuropathy dystrophy and ophthalmoplegia; 5) autosomal recessive progressive external ophthalmoplegia; and 6) autosomal dominant progressive external ophthalmoplegia.\(^12\) Nevertheless, it is increasingly recognized that there is a broad clinical spectrum across these clinical syndromes, in which a patient with **POLG-RD** may present with clinical features that may not fit within a single classically described syndrome.\(^7\)

**POLG** pathogenic variants are the most common cause of mitochondrial epilepsy.\(^8\) Seizures are reported as the first clinical presentation in 50% of all patients with **POLG-RDs**.\(^4\) The most critical factor that triggers epileptic activity is the loss of mtDNA (mtDNA depletion). mtDNA depletion causes loss of the respiratory chain components, which restricts normal energy metabolism, causing a continued neurodegenerative process that interferes with neuronal function. This causes a vicious cycle that ultimately leads to neuronal death and parenchymal necrosis.\(^8\)

EPC is a distinct type of focal motor seizure, first described in 1894.\(^9\) EPC characteristically involves repetitive, sometimes rhythmic, unilateral focal motor twitching of the limbs and/or face, with preservation of consciousness, suggesting an underlying brain lesion.\(^10\) EPC presentation is variable, occurring as single or multiple episodes, and it may be chronic, progressive, or nonprogressive.\(^11\) Several neurologic entities are associated with EPC, such as tuberous sclerosis, Sturge-Weber syndrome, and cortical dysplasia. Metabolic abnormalities such as hypoglycemia; hypoglycemia; hyperuricemia; uremia; brain tumors such as oligodendroglioma, meningioma, and high-grade glioma; autoimmune processes; and infections. Rasmussen encephalitis also has a well-known association with EPC.\(^11,12\)

A comprehensive literature review of 136 patients with epilepsy with **POLG-RDs** has shown that stroke-like changes were the most common imaging findings. The lesions were more commonly located in the occipital lobes. Other structures involved were the parietal, temporal, and frontal lobes; thalamus; basal ganglia; and cerebellum.\(^4\) The age of the patients in this study varied from younger than 30 days to 64 years, with a median of 2 years (first quartile = 0.75 and third quartile = 13.50).\(^4\) Before our study, reports of perirolandic MR imaging signal changes in **POLG-RDs** have only been mentioned in a few isolated case reports, including a 10-month-old child with Alpers-Huttenlocher syndrome who showed multifocal diffusion restriction areas in the left insula, deep gray matter, and
bilateral perirolandic area, which were initially related to a nonspecific metabolic process.
Right pre- and post-central gyri T2 and DWI signal abnormalities were also detected in a 7-month-old girl with POLG-RD who developed focal clonic status epilepticus, mainly of her left arm and occasionally also involving the left leg. Additional case reports described subtle signal changes on FLAIR in a 15-year-old girl with EPC and a 13-month-old child with bilateral perirolandic restricted diffusion with a history of hypotonia, mild motor delay, and EPC.

In this current study of children with POLG-RDs, all patients presented initially with seizures. Most initial brain MR imaging findings were abnormal. A temporal association between seizures and brain MR imaging abnormalities was found, suggesting lesion-related seizures. The primary brain MR imaging abnormalities included unilateral or bilateral signal changes in the perirolandic region, unilateral or bilateral signal changes in the thalamus, or a combination of the 2 in most of our patients. At diagnosis, the perirolandic region or the thalamus was spared in only 2 patients in our cohort, one with unremarkable brain MR imaging findings and the other in whom diffuse white matter with restricted diffusion was noted (Fig 3).

The perirolandic sign was common and present in most patients at diagnosis. Moreover, the sign was seen in most patients during the course of the disease because half of the patients who did not have the sign initially presented with the sign later. The presence of perirolandic signal changes would reflect neuronal death following acute energy failure in a metabolically demanding region (primary motor and sensory cortex) within the dominant hemisphere. Further studies are needed to confirm this hypothesis.

The MR imaging appearance of the perirolandic sign was varied. Signal changes involved both the pre- and postcentral gyri, more commonly in the precentral gyri. The conspicuity of the perirolandic sign was also variable (Fig 1). In the more notable cases, signal changes were ribbon-like following the course of the gyri, which were readily detectable as T2 hyperintensities and restricted diffusion. DWI was the most sensitive MR imaging sequence to detect signal changes and, therefore, should always be included in the protocol and carefully evaluated when a case of POLG-RD is suspected.

Thalamic signal changes were also frequent at the time of diagnosis and on follow-up imaging. Unilateral or bilateral thalamic involvement was identified in most patients during the onset of their disease. In our cohort, there was no new thalamic involvement at follow-up. On follow-up imaging, thalamic changes had different outcomes: complete resolution, progression from unilateral to bilateral involvement, progression accompanied by volume loss, and fluctuation with periods of an almost-complete resolution and frank progression. Thalamic signal changes were variably detected on DWI, T2WI, FLAIR, or ASL.

Volume loss was detected almost always on follow-up imaging. Volume loss varied from mild to severe, showing rapid evolution in most cases. Most important, volume loss showed no lobar predominance in most patients. However, in severe cases, the occipital lobes were affected asymmetrically and with encephalomalacic changes. Signal changes were detected virtually in any part of the brain, except the medulla. The spinal cord was not part of the scope of this study.

The occipital lobe was rarely affected at diagnosis, though frequently involved on follow-up imaging. Advanced MR imaging sequences such as MR spectroscopy and ASL, albeit not...
being acquired in all the patients, were useful to demonstrate increased lactate peaks and areas of localized hyperperfusion, respectively. An increased lactate peak is an expected finding in primary mitochondrial disorders and has already been described in patients with POLG-RDs, reflecting mitochondrial dysfunction.\(^8\) ASL perfusion has mostly been described in patients with mitochondrial encephalomyopathy with lactic acidosi and stroke-like episodes (MELAS), but rarely in POLG-RDs. Increased ASL perfusion in MELAS preceded signal changes on conventional MR imaging in 3 patients, indicating that ASL imaging has the potential for predicting the emergence of stroke-like lesions.\(^17\) In our cohort, increased ASL perfusion was coincident with either T2WI, FLAIR, or DWI signal changes and may have been related to seizure activity. ASL, however, was useful to indicate areas that would undergo further volume loss in some of our patients (Fig 5). The complete role of ASL perfusion in patients with POLG-RDs needs further investigation.

Valproic acid is an effective antiepileptic drug to manage epilepsy. However, it is strictly contraindicated to treat seizures in patients with POLG-RD due to its potential to cause fulminant hepatotoxicity.\(^9\) The best diagnostic test to prevent valproic acid–induced liver toxicity is POLG gene testing before the use of this medication.\(^18\) The presence of a unilateral or bilateral periorl and occipital increased perfusion is seen in C (open arrows) and in the bilateral cerebellar hemispheres (open arrows) in D. Increased ASL perfusion is seen in the left thalamus in E (solid arrow) and in the left parietal and occipital lobes in E and F (open arrows).

MR imaging findings in our cohort are not pathognomonic for POLG-RD, and the differential diagnosis includes postictal changes and stroke-like episodes similar to MELAS. At this time, we can only speculate about their nature, and further laboratory, genetic, and imaging correlation are necessary to establish their pathogenesis. Bilateral periorl and occipital increased perfusion is seen in C (open arrows) and in the bilateral cerebellar hemispheres (open arrows) in D. Increased ASL perfusion is seen in the left thalamus in E (solid arrow) and in the left parietal and occipital lobes in E and F (open arrows).
and family meetings hosted by United Mitochondrial Disease Foundation; European Society of Human Genetics; Society for the Study of Inborn Errors of Metabolism; Society of Inherited Metabolic Disease; Hebrew University; Asian Society Of Inborn Errors of Metabolism; Wellcome Genome Campus; Cold Spring Harbor Laboratory. Amy Goldstein—UNRELATED: Board Membership: United Mitochondrial Disease Foundation. Comments: Scientific and Medical Advisory Board. *Money paid to the institution.

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