

Case report



Case Report of a de Novo Mutation of PCDH19 in Saudi Family Limited to Females

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Abstract

Up to date more than 60 different mutations in PCDH19 have been identified. Most of PCDH19 gene is located in Xq22 and produces nonclustered delta protocadherin. This disorder primarily manifests in heterozygote females due to random X chromosome inactivation leading to somatic mosaicism and abnormal cellular interference between cells with and without delta-protocadherin., but we a heterozygous nucleotide mutation causing amino acid 561 to change from Pro to Ser (p.Pro561Ser). This mutation was de novo, and this alteration was not found in her parents. PCDH19-related epilepsy is a distinct childhood-onset epilepsy syndrome characterized by brief clusters of febrile and afebrile seizures with onset primarily before the age of three years, cognitive impairment, autistic traits, and behavioral abnormalities. We describe the features of a de novo mutation in 3 sibling, presented with early onset of seizure, two of them were controlled and wean off medication was at age of six year and her sister at age of 10 year .The youngest sister still partially controlled on medication, she had seizure only during febrile illness.

Keywords: Genetic; PCDH19; focal seizure; sibling; anti-epileptic medications

Case report

Here we are describing three sisters (13 year-8 year- 5-year-old girl) who suffered from repetitive seizures. The oldest sister 13-year-old girl developed a febrile seizure at the age of 14 months characterized by generalized loss of tone with impaired awareness lasted for few minutes. Admitted initially as febrile status for investigation suspected post viral infection to rule out CNS infection. Her examination was normal neurological exam, no focal deficit and her systemic review was within normal, no sore throat or ear infection was noted. Blood work up was within normal (blood count , Chemistry, renal function) was all normal , apart from mild leukocytosis (white blood cell 12.4), cerebral spinal fluid (CSF) analysis and culture was negative , virology culture was negative. Patient was discharge after few days in stable condition and no anti-epileptic medication was started. Later at age of 2 year

and 3 months, she started to have afebrile seizures. The seizures occurred 3-4 times per week, similar to her previous description generalized tonic clonic seizure with no focality lasting 3-5 minute followed by post ictal sleep or drowsy for few hours till she is back to her baseline. Her EEG using the international 10-20 system showed generalized spike and wave. MRI brain was normal. She was diagnosed to have G.epilepsy, started on Topiramate; however, her seizures were not responsive. The longest period of remission was just 3-4 weeks. Then levetiracetam16 was introduced, later showed improvement in-term of frequency and duration (Valproic acid was not recommended but parents refuse concern about the side effect). EEG was repeated showed normal recording .She was born at full-term with a birth weight of 3.5 kg. She had normal early developmental milestones. She was the first child of unrelated parents. There was no family history of epilepsy or intellectual disability. The patients' family pedigree is shown in Figure 1.

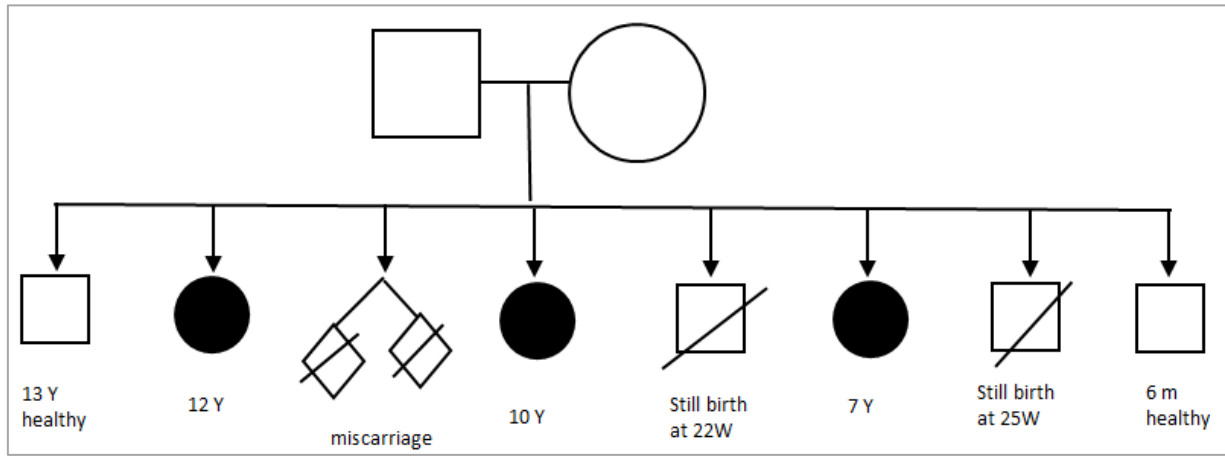


Figure 1

The second sister, an eight-year-old girl, was reported to have repetitive seizures starting at the age of 14 months. She was born at full-term without birth-related distress or dysmorphism. Perinatal history and pregnancy were both unremarkable. Her seizure started as febrile, described as a tonic seizure following by screaming and a frightened face, including loss of contact, stiff limbs, purple lips, and chewing movements. These episodes lasted for about 30 sec. Later, the seizures appeared without fever with similar semiology. EEG showed using the international 10-20 system, generalized spikes and wave higher frontal bilateral, in some epochs showed few spikes at posterior temporal at P3- T5, the last EEG showed normal recording, and an MRI brain scan is unremarkable. Levetiracetam (LEV) [16] was started as her older sister; the patient experienced reduced attack frequency, she was on Levetiracetam for more than 2 year and at age of 5 year start to wean off medication, patient had 2 attack of generalized seizure after few months from stopping medication, Levetiracetam was continue till age of 8 year, was seizure free then start to wean off currently she is 9 year and 6 months. The onset of epilepsy seizures was accompanied by no significant developmental delay. The patient’s intelligence level was not assessed by appropriate scale, whether before or after her onset. But during the follow-up period, her mother reported overall good school performance. Third sister is a 5-year-old girl. She was born at full-term via with no distress or dysmorphic features. The perinatal history and neonatal course were all unremarkable. Her first seizure displaying a right focal tonic seizure was noted during a febrile illness at the age of 9 months. The EEG done using the

international 10-20 system was showing normal posterior dominant rhythm and frequent spike and wave seen at posterior head at P3-O1 and P4-O4 with lesser extent to T6, the last reading was repeated after 8 months from initiation of medication was normal recording and no epileptiform activity. Then the brain computed tomography (CT) done reported normal. She developed a cluster of febrile seizures by 19 months of age, lasting for approximately 10 s each time occurred around six times in 36 h. By 31 months of age, a second cluster appeared triggered by fever, her seizure semiology change to sudden behavior arrest. The brain MRI was also normal at this onset. Her psychomotor development was unremarkable. She received multiple antiepileptic treatments, including sodium valproate, Topiramate, phenobarbitone, and lamotrigine (started for few weeks develop skin rash, medication wean off). The longest period of remission was just two months. Her antiepileptic regimen was adjusted multiple times. Her seizure was controlled after being given carbamazepine, levetiracetam. She is seizure free for more than 1 year and 5 months. Whole-exome sequencing result showed a missense mutation in the PCDH19 gene: a cytosine to guanine (C > G) nucleotide change at position 1031 (c.1031C > G), a heterozygous nucleotide mutation causing amino acid 561 to change from Pro to Ser (p.Pro561Ser). This mutation was de novo, and this alteration was not found in her parents. Three mutations of PCDH19-related epilepsy were identified in three children within the same family. The clinical data and characteristics of the three patients carrying PCDH19 variations are described in Table 1.

Table 1

Patient	Age	Gender	Age at seizure onset	Seizure type at onset	Other seizure type	EEG	Brain Image	Development before seizure onset
1	5y	female	9 M	Right focal	Generalized, focal Eye gaze to the left with GTC	Focal, posterior head	unremarkable	normal
2	8y	Female	14 M	Screaming and GTC	non	GSW	unremarkable	normal
3	13y	Female	14 M	Tonic seizure	Eye gaze to the left then GTC	GSW	unremarkable	normal

Table 2

Patient	Medication tried	Current medication	Seizure outcome	Current cognitive status
1	Topiramate velprate phenobarbitone lamotrigin	Carbamazepine, levetiracetam	Seizing on ASMs	Normal
2	levetiracetam		One year off meds, No seizure	Normal
3	Topiramate	levetiracetam	Controlled, seizure free, off medication	Difficulty in memorizing

Discussion

We had notice in all the 3 cases, each patient had different semiology of seizure and their response of anti-seizure medication was variable in spite all of them had same gene mutation and similar early presentation (febrile seizure). But there was clear good response of medication and the outcome was also good in 2 of the siblings although the youngest she showing good response apart from the intermittent breakthrough of seizure that provoked with fever. There was no clear description was mentioned in the other literature and if there is clear variability of seizure presentation in sibling. In literature, around 292 cases were reported in different studies [1]. Approximately 150 mutations of PCDH19 have been discovered, with more than half affecting extracellular adhesive domains by missense mutations as found in our patients [2]. Here, the findings from our study are compared with those of previously reported cases [1]. In previous reports, PCDH19 cases share the following features: age at the onset before three years, seizures semiology described as earlier fearful screaming, occurred in the cluster and cognitive and behavior disturbances [3,4]. Similarly, in our three patients, the seizures onset in the first year of life for one patient and after the first year of life for the other two, with an average age of onset of 12 months (9–14 months). All patients developed generalized seizures or focal seizures evolve to secondary generalization. Seizures occurred in clusters with all subjects, and patient?? Had confirmed epileptic status. All three patients had seizures associated with fever sensitivity at the onset. Patient 3 had recurrent seizures despite being treated with multiple anti-epileptic drugs, suggesting that PCDH-19 related seizures had variations and were difficult to control to a certain degree [5]. Interictal EEG was normal in all patients, similar to what had been described in the literature. Moreover, Fast activity over a localized brain region was also described as an interictal finding. The majority of described Ictal EEG in literature is focal onset being frontotemporal most common, though generalized seizure onset has been reported [1]. Brain MRI had no particular findings in all three patients, as reported in most previous cases. However, a minority of reported cases were found to have cortical malformation and hippocampal sclerosis [6]. Psychiatric symptoms were detected among all the patients. However, patient 1 has memorizing difficulty and patient 3 developed intellectual deterioration and behavioral disturbance. Out of previously reported cases, 213 patients of 271 patients had a psychiatric illness. Autism and ADHD are mostly reported [7,8]. Regardless of the variation in clinical manifestations of PCDH19-related epilepsy, the disease shares some common characteristics, particularly the age of onset at infancy and the seizure semiology as generalized seizures and cluster seizures [9]. The most common seizure types reported are generalized tonic seizures, tonic-clonic seizures, or focal seizures with secondary generalization appearing in clusters with or without fever. Other seizure types are rare such as myoclonic seizures, atonic seizures, and atypical absences. Cognitive development vary from normal to severe intellectual impairment. In addition, Patients may or may not demonstrate psychiatric symptoms and behavioral disturbance [10,11]. Of the three patients, three genetic alterations were the same exact location suggesting inherited from their asymptomatic father. (Carrier status??) PCDH19 mutations in most patients are sporadic and de novo, while some are of paternal inheritance. This makes it challenging to recognize the pattern of inheritance [12,13]

In one patient, treatment with levetiracetam was reported as effective. In the other patients, treatment with oral polypharmacy,

clonazepam (CZP), and phenobarbital (PB) also was reported as effective [14,15].

Conclusion

Genetic testing play a good role in diagnosis of familial epilepsy but with minimal knowledge of medication effect in those type of mutation. In our report the choice of antiepileptic medication is limited but there was response on levetiracetam as adjunctive medication but again the efficacy of medication response and better choice need base evidence.

Conflict of interest

None

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