


Cognitive impairment in children with *CACNA1A* mutations

VERONIQUE HUMBERTCLAUDE^{1,*} | FLORENCE RIAN^{2,3,*} | BENJAMIN KRAMS⁴ | VALERIE ZIMMERMANN⁴ |
NICOLAS NAGOT⁵ | DANIEL ANNEQUIN⁶ | BERNARD ECHENNE⁴ | ELISABETH TOURNIER-LASSERVE^{2,3} |
AGATHE ROUBERTIE^{4,7}  | EPISODIC SYNDROME CONSORTIUM[†]

1 Service de Médecine Psychologique Enfants et Adolescents, CHU Saint Eloi, Montpellier; **2** Laboratoire de Génétique, Groupe hospitalier Lariboisière-Fernand Widal AP-HP, Paris; **3** INSERM UMR-S740, Université Paris 7 Denis Diderot, Paris; **4** Département de Neuropédiatrie, CHU Gui de Chauliac, Montpellier; **5** Centre d'Investigation Clinique, CHU Montpellier, Montpellier; **6** Centre de la Migraine de l'Enfant, Hôpital Trousseau, APHP, Paris; **7** INSERM U 1051, Institut des Neurosciences de Montpellier, Montpellier, France.

Correspondence to Agathe Roubertie, Département de Neuropédiatrie, CHU Gui de Chauliac, 80 Avenue Fliche, 34295 Montpellier, France. E-mail: a-roubertie@chu-montpellier.fr

*These authors contributed equally to the manuscript.

†Members of the Episodic Syndrome Consortium are listed in Appendix S1 (online supporting information).

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ABBREVIATIONS

BPT	Benign paroxysmal torticollis
BPV	Benign paroxysmal vertigo
BTU	Benign paroxysmal tonic upgaze

AIM To describe the clinico-radiological phenotype of children with a *CACNA1A* mutation and to precisely evaluate their learning ability and cognitive status.

METHOD Children between the ages of 3 and 18 years harboring a pathogenic *CACNA1A* mutation associated with episodic ataxia, hemiplegic migraine, benign paroxysmal torticollis, benign paroxysmal vertigo, or benign paroxysmal tonic upgaze, were enrolled in this cross-sectional study. Data concerning psychomotor development, academic performance, educational management, clinical examination at inclusion, and brain imaging were collected. Cognitive assessment was performed using age-standardized scales.

RESULTS Eighteen patients (nine males, nine females; mean age at inclusion: 11y 7mo [SD 4y 5mo; range 3y–17y 11mo]) from 14 families were enrolled. Eleven patients displayed the coexistence or consecutive occurrence of more than one type of episodic event. Nine patients exhibited abnormal neurological examination at inclusion. Brain magnetic resonance imaging (MRI) showed cerebellar atrophy in five patients. Psychomotor development was delayed in nine patients and academic difficulties were reported by the parents in 15 patients; nine patients were in special education. Impairment of intellectual function was assessed in six of the 12 patients with interpretable Full-scale IQ scores and was more frequent when cerebellar atrophy was present on MRI.

INTERPRETATION Cognitive impairment is commonly associated with *CACNA1A* mutations. We suggest that *CACNA1A*-associated phenotype should be considered a neurodevelopmental disorder.

The *CACNA1A* gene, located on chromosome 19p13, encodes the pore-forming Cav2.1 (alpha 1A) subunit of P/Q-type Ca²⁺ channels, which plays a major role in neurotransmitter release throughout the nervous system, especially in the cerebellar Purkinje cells.¹ Triplet repeat mutations in the *CACNA1A* gene are associated with a homogeneous phenotype of late onset-spinocerebellar ataxia type 6 (OMIM 183086).² Heterozygous point mutations or deletions of the *CACNA1A* gene are associated with a heterogeneous spectrum of episodic and/or permanent neurological manifestations. More than 20 years ago, mutations in the *CACNA1A* gene were identified in patients with hemiplegic migraine (OMIM 141500) and in patients with episodic ataxia type 2 (OMIM 108500).³ More recently, single case reports or short case series showed the involvement of *CACNA1A* mutations in various types of episodic neurological manifestations with or without hemiplegic migraine or

episodic ataxia type 2, episodic syndrome that may be associated with migraine (benign paroxysmal torticollis [BPT]), benign paroxysmal vertigo (BPV), benign paroxysmal tonic upgaze (BTU), epileptic seizures, or dyskinesia.^{4–10} Ataxia, with or without cerebellar atrophy, has been reported in adults, and in single case reports of children with *CACNA1A* mutations.^{6,11–16} Among the chronic neurological manifestations associated with heterozygous *CACNA1A* mutations, cognitive dysfunction has been suggested in recent years, including developmental delay, learning difficulties, autism, attention-deficit/hyperactivity disorder, intellectual impairment, and epileptic or non-epileptic encephalopathy.^{6,17–23} Until now, the cognitive profile of children with *CACNA1A* mutations has not been studied systematically in a series of patients.

This study aimed to describe the clinico-radiological phenotype of children with *CACNA1A* mutations, with

special focus on psychomotor development and cognitive abilities, and to determine the clinical and genetic factors associated with cognitive dysfunction.

METHOD

Between July 2011 and February 2016, 18 patients were recruited through the Childhood Episodic Syndromes Study, a large, multicentre, research project on episodic syndromes in children, part of whose results have recently been published.²⁴ Patients were included in the study if they were: (1) between 3 and 18 years old; (2) had a pathogenic *CACNA1A* mutation; and (3) they (or their parents) agreed to undergo clinical and neurocognitive evaluations. Thirteen patients were tested for *CACNA1A* mutations before the study and were included for clinical evaluation and five had both molecular and clinical evaluation during the multicentre research project (Figure S1, online supporting information). Eleven University Hospitals in France were involved in this cross-sectional study.

The study was carried out in accordance with the Declaration of Helsinki and approved by the Sud-Méditerranée ethical committee. Written informed consent for clinical data collection and genetic studies was obtained from the patients' legal representative.

Clinical assessment

The clinical data were collected by seven delegated participating clinicians using a structured questionnaire specifically designed to gather information concerning perinatal events, psychomotor development, familial history, and neurological examination at inclusion. The following data were collected concerning the occurrence of any type of paroxysmal neurological events: type of event (ataxia, BTU, paroxysmal torticollis, vertigo, other forms of movement disorders, hemiplegic or non-hemiplegic migraine, epileptic seizures), age at onset, duration of the episodes, treatments, and outcome at inclusion. Episodic events were diagnosed according to International Headache Society criteria^{25–27} or diagnosis criteria accepted in the literature, this is summarized in Table SI (online supporting information). Magnetic resonance imaging (MRI) data were reviewed by an experienced neuroradiologist (NL, see Appendix S1) and a paediatric neurologist (AR).

Information concerning rehabilitation (speech, occupational, or psychomotor therapy) for difficulties in academic performance, specific assistance in regular school (services provided by the Ministry of Education in France such as computer use or human assistance), and education in special schools were collected from the medical files and during familial interview at inclusion.

Standardized cognitive scales were administered according to the child's age: the Wechsler Preschool and Primary Scale of Intelligence, Third Edition for children under 7 years, the Wechsler Intelligence Scale for Children, Fourth Edition for children between the ages of 5 and 16 years 11 months, or the Wechsler Adult Intelligence Scale, Third Edition for children older than 16 years.

What this paper adds

- Cognitive disabilities and academic difficulties are common in children with *CACNA1A* mutations associated with episodic syndromes.
- Cognitive function ranges from normal to moderate intellectual disorder in wheelchair-dependent children.
- Patients with vermian atrophy are at a higher risk of cognitive impairment.

Genetic assessment

Screening of the *CACNA1A* gene was performed by sequencing and quantitative multiplex polymerase chain reaction of short fluorescent fragments analysis. The mutations present in the patients were considered as deleterious according to the following criteria: they were mutations leading to a stop codon or they had already been reported in families experiencing hemiplegic migraine or episodic ataxia (p.T666M, p.G677R, p.R1352Q); they appeared de novo (p.Y1706_I1709del) or were: (1) absent in data bases; (2) prediction tools (polyphen 2, Scale-Invariant Feature Transform, Align Grantham Variation and Grantham Deviation, and mutation taster) gave results in favor of a deleterious effect on the gene; (3) they were localized in a P-loop domain critically implicated in channel function; (4) the patient's phenotype was typical of a *CACNA1A* linked disease; and (5) they cosegregated with the disease in the tested family members (p.C1417W, p.T1458M).

Statistical analysis

Patients' characteristics were presented using median and interquartile range (IQR), mean and standard deviation (SD), or range for continuous variables, frequencies, and proportions for categorical variables.

Groups of patients (defined by deficit in intellectual function and cerebellar atrophy) were compared using Student's *t* or Wilcoxon–Mann–Whitney *U* tests, depending on the normality of the distributions tested with Shapiro–Wilk test, for continuous variables and Fisher's exact test for categorical ones.

Statistical bilateral significance threshold was set at 5%. Statistical analyses were performed using SAS, version 9.1 (SAS Institute, Cary, NC, USA).

RESULTS

Results are reported in Table SII (online supporting information).

Population

Eighteen patients (nine males, nine females) from 14 families were enrolled, including three sib-pairs (patients 4 and 5, 7 and 8, 12 and 13) and one pair of first cousins (patients 15 and 16). The past medical history of the patients was unremarkable. The mean age at inclusion was 11 years 7 months (SD 4y 5mo; range 3y–17y 11mo). Patients 7, 8, 12, and 18 have been reported elsewhere.^{4,12,28}

Fourteen different mutations were identified: eight mutations leading to a stop codon, one splice site mutation, and five missense mutations (three had already been reported and two were novel).

Episodic events

The episodic events considered in the patients were episodic ataxia, hemiplegic migraine, BTU, BPT, and BPV. Details are shown in Table SIII (online supporting information). Seven patients exhibited only one type of episodic event at last follow-up: episodic ataxia occurred in three patients, hemiplegic migraine in two patients, and BTU and BPT in one patient each. Eleven patients presented a combination of various episodic events. Episodic ataxia or hemiplegic migraine occurred in 15 of 18 patients and were preceded by infantile episodic events (BTU, BPT, or BPV) in five out of 15 patients. Altogether, episodic ataxia occurred in 11 patients, hemiplegic migraine in five patients, BTU in four patients, BPT in three patients, and BVP in one patient. Two patients presented with seizures and myoclonus was reported in one patient.

The first episodic manifestation occurred at a median age of 24 months (IQR 8mo–5y 7mo; range 2mo–14y). BTU, BPT, and BPV were the earliest manifestations in eight patients, with an onset before 4 years of age in all patients, and in the first year of life in six out of eight patients. The number of episodes varied widely. In the eight patients with BTU, BPT, or BPV, the episodic manifestations were considered cured at the time of inclusion, with a median duration of disease of 16 months (IQR 14–23mo; range 13mo–6y 3mo).

Episodic ataxia was the first manifestation in six patients and followed another type of episodic event in five patients. The median age at onset was 36 months (IQR 18mo–9y 6mo; range 1mo–13y). The frequency of episodes varied widely among the patients, ranging from some per year up to almost daily episodes (600 per year in one patient). Hemiplegic migraine occurred as a first phenotype in four patients (patients 15–18), and followed another phenotype in one patient (patient 9). The median age of the start of hemiplegic migraine was 9 years (IQR 7–10y; range 4–14y). The number of episodes during the first year was less or equal to 12 per year in four out of five patients with hemiplegic migraine and varied widely at follow-up. Episodic ataxia and hemiplegic migraine were considered ongoing manifestations at the time of enrollment in all the patients.

Information concerning treatment of the episodic events was collected in 10 patients with various phenotypes: episodic ataxia (six patients), hemiplegic migraine (three patients), migraine (three patients) (Table SIV, online supporting information). Acetazolamide was effective in one of five patients with episodic ataxia and in the two patients with hemiplegic migraine. Lamotrigine resulted in no effect (one of three patients) or aggravation (two of three patients) with episodic ataxia.

Neuropsychological assessment

Data concerning psychomotor development, neurological examination, academic performance, and cognitive assessment are detailed in Table SII and summarized in Table I.

Development and neurological examination at enrollment

Nine patients presented with motor and/or language developmental delay. At the time of enrollment, neurological examination was abnormal in nine patients. Nystagmus was observed in seven patients, associated with cerebellar signs or unsteadiness in three patients, including one severely affected patient with intellectual disability, dyskinesia, and inability to walk alone (patient 17). Cerebellar signs with hypotonia or isolated hypotonia were disclosed in two other patients. Three patients exhibited early-onset non-progressive cerebellar signs, mimicking congenital ataxia (patients 4, 14, and 17). No other neurological symptoms were recorded (including autistic features).

Academic performance and rehabilitation therapy

Significant difficulties in academic performance were reported in 15 patients. Nine patients attended special schools; among the nine patients who were in a regular school, four benefited from specific learning assistance. Thirteen among the 18 patients benefited from rehabilitation (speech and/or occupational, and/or fine motor rehabilitation) therapy.

Cognitive assessment

IQ was evaluated in 16 patients: two with the Wechsler Preschool and Primary Scale of Intelligence, Third Edition, 13

Table I: Psychomotor development, neurological examination, learning disabilities, educational management, and cognitive assessment of the patients

		Number of affected patients/number of patients with known data
Neuropsychological assessment		
Neurological examination at inclusion	Abnormal	9/18
	Nystagmus	4/18
	Nystagmus+cerebellar signs or unsteadiness	3/18
	Cerebellar signs	1/18
Psychomotor delay	Hypotonia	1/18
	Motor delay	6/18
Schooling	Language delay	6/16
	Difficulties in academic performance according to families	15/18
Rehabilitation	Special education school	9/18
	Regular school with specific learning assistance	5/18
	Regular school without specific learning assistance	4/18
	Total	13/18
Full-scale IQ	Speech therapy	13/18
	Occupational therapy	3/18
	Psychomotricity	5/18
	<70	6/12
	70–79	4/12
	80–119	2/12
	≥120	0/12

with the Wechsler Intelligence Scale for Children, Fourth Edition, and one patient with the Wechsler Adult Intelligence Scale, Third Edition (Table SII). Two families completed the study but refused cognitive evaluation. Full-scale IQ was not interpretable in four patients because of excessive discrepancy among the subscales; therefore, Full-scale IQ scores were available in 12 patients. Moderate intellectual disability (Full-scale IQ<50) was present in four out of 12 patients, mild intellectual disability (50–69) in two out of 12 patients; four out of 12 patients had borderline intellectual disability (70–79). Full-scale IQ was in the average range (80–119) in two out of 12 patients, and no patient had an IQ in the superior range (≥ 120 ; Fig. 1). Detailed analysis of IQ scores in the whole group of patients did not reveal a specific pattern of cognitive impairment.

Brain imaging

Twenty-eight MRI scans were available for the 18 patients. Atrophy of the superior part of the cerebellar vermis was detected in five patients (Fig. 2). The atrophy was present at 4 years and 1 month (patient 14), 11 years and 9 months (patient 9, for whom it was absent at 4y 11mo), 12 years and 9 months (patient 10), 15 years 5 months (patient 5), and 15 years 9 months (patient 17). Three patients exhibited episodic ataxia and two had hemiplegic migraine, preceded by BPV in one patient. Of the five patients, four had an abnormal neurological examination: one had isolated hypotonia and three had a nystagmus, associated with cerebellar signs for two of them. Four of the five patients exhibited a deficit in intellectual function, and patient 9 did not undergo IQ assessment.

Family history

The occurrence of episodic manifestation appeared to be sporadic in six patients. The parents of patient 1 and mother of patient 10 did not have the mutation that was found in their child. The father of patient 9 was an asymptomatic carrier. The other parents were not available for testing.

The eight other patients were familial forms. The mutation was present in a total of 20 additional relatives (Fig. 3). Symptomatic family members of patients 15 and 16 with mutations exhibited the same phenotype (severe attacks of hemiplegic migraine without cognitive dysfunction). In the other seven families, intrafamilial phenotype variability was observed (Table SII). In particular, various associations of BTU, BPT, episodic ataxia, migraine, and cognitive dysfunction were observed in six members of the family affected by the stop mutation c.2206C>T (two sisters, [patients 7 and 8], their mother, a cousin, and a maternal uncle).⁴ BTU and episodic ataxia were associated in five of the 18 families, successively in the same patient (patients 1, 2, 3, and the cousin of patients 7 and 8) or in the same family (two sisters, patients 4 and 5).

Correlations

Deficit in intellectual function (Full-scale IQ<70 in six of 12 patients) tended to be correlated with cerebellar atrophy on brain MRI (four of six patients with Full-scale IQ<70 vs zero of six patients with Full-scale IQ ≥ 70 , $p=0.06$). No significant relation could be established between a deficit in intellectual function and sex, young age at onset of the first episodic event, type of episodic event, and occurrence

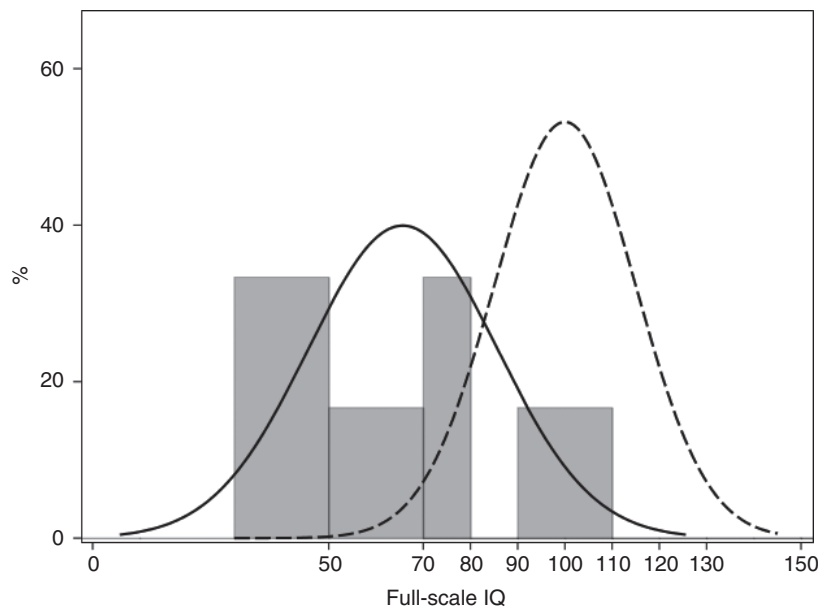


Figure 1: Distribution of the Full-scale IQ score (16 patients). This figure demonstrates the increased incidence of patients with low (<70) or border line (<80) IQ among patients with a *CACNA1A* mutation. Dotted line, general population; solid line, patients with a *CACNA1A* mutation; grey bars, number of patients. [Correction added on 13 June 2019 after first publication: the figure caption has been corrected.]

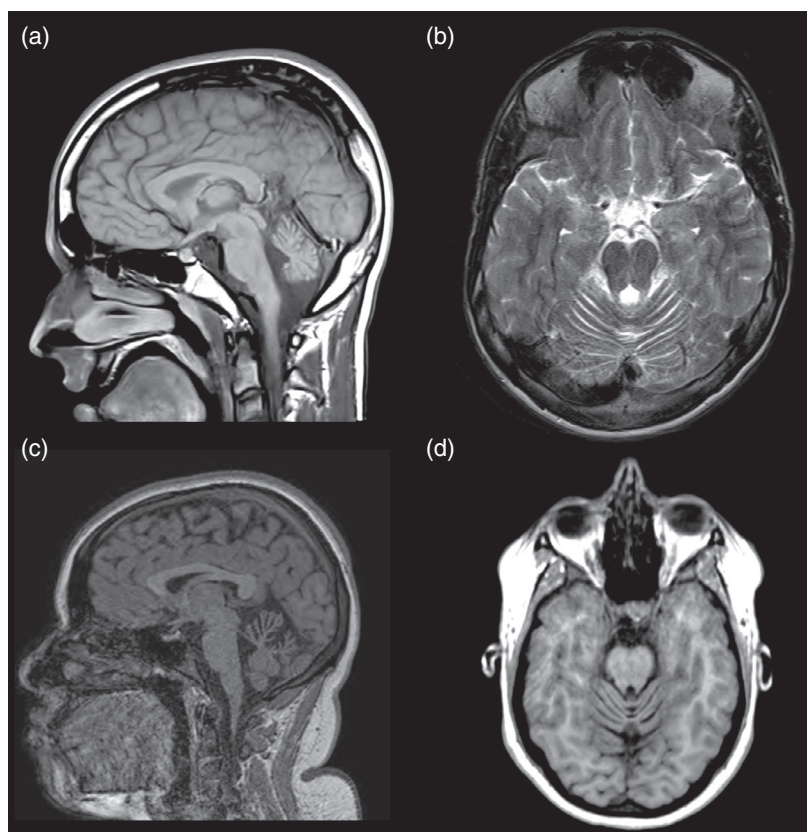


Figure 2: Brain imaging illustrating the vermian atrophy in (a,b) patient 17 and (c,d) patient 9. (a) Sagittal T1 (b) and axial T2 images, cerebellar atrophy mainly involving the superior vermis; (c) sagittal (d) and axial T1 images, cerebellar atrophy involving the superior and posterior vermis with enlarged primary fissure.

of various types of episodic events, abnormal neurological examination, or type of mutation.

The five patients with cerebellar atrophy exhibited motor delay more frequently than patients with normal brain MRI (four of five patients vs two of 13 patients, $p=0.021$). The proportion of stop mutations was lower in patients with cerebellar atrophy (one of five patients vs 10 of 13 patients, $p=0.047$). Patients with abnormal MRI tended to be older (mean age [SD] 14y [2y 11mo] vs 10y 8mo [4y 7mo], $p=0.15$). No significant association could be established between cerebellar atrophy and sex, young age at onset of the first episodic event, abnormal neurological examination, and type of episodic events or occurrence of various types of episodic events.

DISCUSSION

This study describes the clinical and radiological phenotype of 18 children from 14 families with *CACNA1A* mutations. The patients had a detailed clinical description and benefited from a standardized neurological examination and evaluation of psychomotor development and academic performance, and a cognitive assessment. They also underwent neuroimaging. Our results confirm a diversity of episodic attacks in the patients and a frequent association of

neurological abnormalities, in particular, an abnormal cognitive profile.

Precise cognitive evaluations are the main contribution of our study. Recently, Indelicato et al.²³ reported cognitive and behavioral features in a series of patients with *CACNA1A* mutations. However, the authors performed a retrospective analysis of data extracted from a database initially developed for another objective, therefore the protocol assessment of cognitive function was not uniform. Our study reports the first series performed with prospective and standardized neuropsychological assessment. It shows that cognitive disabilities are associated with *CACNA1A* mutations in half of the children with available data, and that cognitive dysfunction is heterogeneous, ranging from a moderate disorder of intellectual development in wheelchair dependent patients to mild difficulties at school. The analysis of the different subtests of the Full-scale IQ testing did not disclose a specific cognitive profile; several domains of cognitive function were impaired in our patients (data not shown). Autistic symptoms that have been previously reported in two families with episodic ataxia type 2 were not detected in our patients.²¹ Vermian atrophy was linked to a higher risk of cognitive dysfunction.

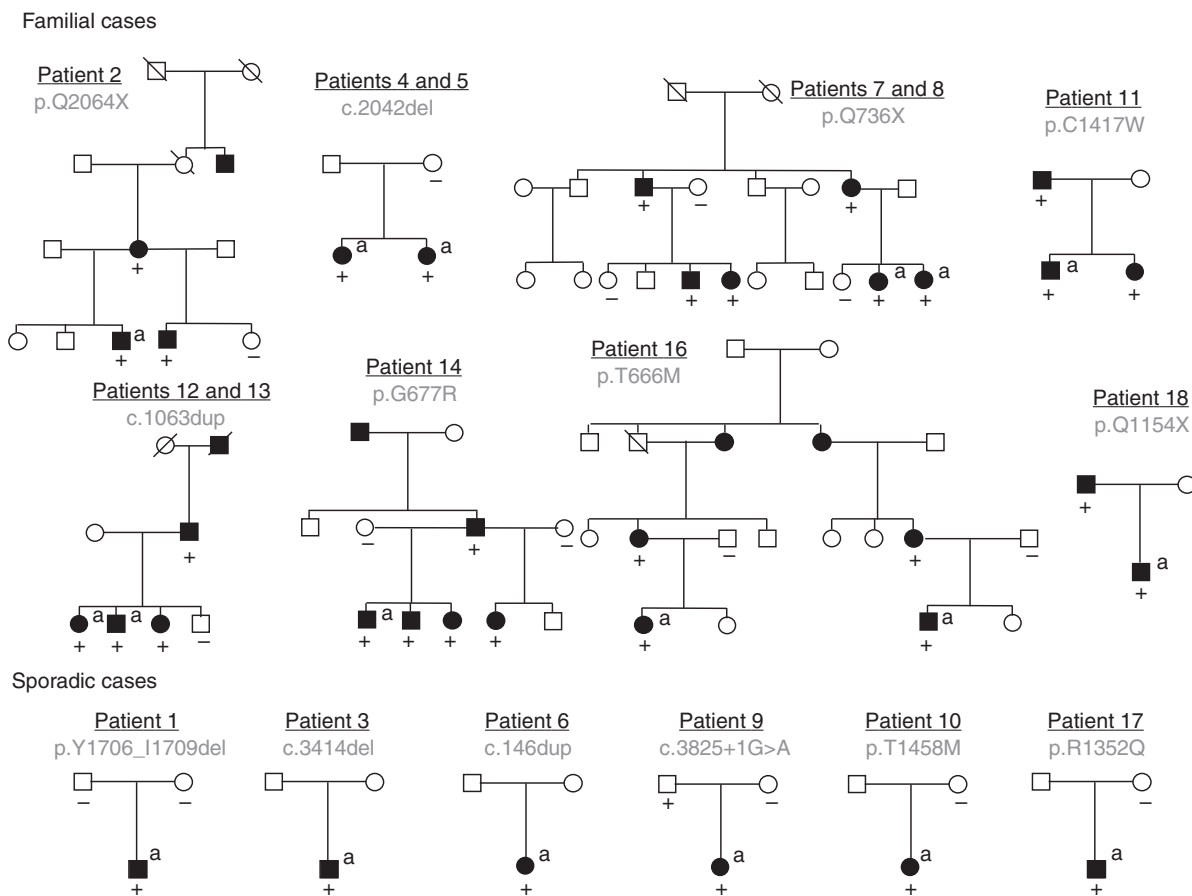


Figure 3: Pedigrees of the patients. Squares represent males and circles represent females. Black symbols represent individuals with episodic symptoms (hemiplegic migraine, episodic ataxia, tonic upgaze, or paroxysmic vertigo). White symbols indicate unaffected individuals, and symbols with a diagonal line represent deceased individuals. ^aPatients included in this study. +, a mutated allele; -, normal allele.

The small size of our patient sample represents a clear limitation if this work, especially regarding establishing a correlation without considering the possible within-family correlations. Given the young age of some patients at inclusion, the results concerning academic difficulties and rehabilitation might underestimate these difficulties and the need for rehabilitation in patients with *CACNA1A* mutations. Recently, pharmaco-sensitive, but also devastating drug resistant epilepsy has been reported in patients with a *CACNA1A* mutation;^{19,29} however, the prevalence of epilepsy in patients with a *CACNA1A* mutation is still unknown. Given that our patients were recruited from the Childhood Episodic Syndromes Study and the database from Lariboisiere laboratory, which researches migraine and syndromes associated with migraine, patients with epilepsy and a *CACNA1A* mutation might be under represented in our series. In fact, only two patients from our series exhibited epilepsy (absence epilepsy/myoclonus), therefore, our study cannot provide further clues to the issue of the role of epilepsy in cognitive impairment in patients with *CACNA1A* mutations.

Our study shows that the phenotype of the patients with *CACNA1A* mutations is highly variable in terms of diversity of clinical manifestations, but also in terms of an association of several types of symptoms, as summarized in Figure S2 (online supporting information). Symptoms include non-epileptic manifestations (ataxia, torticollis, vertigo, tonic upward gaze, migraine), and epileptic events (absence seizures, myoclonus). A combination of several types of episodic syndromes was present in two out of three of our patients, either concomitant or evolving from one type to another. Permanent neurological abnormalities were also present in half of the patients at inclusion, consisting of variable degrees of cerebellar dysfunction (nystagmus, ataxia, hypotonia), with or without vermian atrophy. Our study confirms that infantile-onset episodic syndromes, such as BTU or BPT, may be precursor of *CACNA1A*-related episodic ataxia or hemiplegic migraine,⁸⁻¹⁰ since in one-third of the patients, occurrence of episodic ataxia or hemiplegic migraine was preceded by BTU, BPT, or BPV.

Our study clearly highlights the wide phenotypic spectrum associated with *CACNA1A* mutations, regarding age

at onset of the symptoms, cognitive profile (ranging from typical to moderate disability), episodic syndromes (diversity of the syndromes), and neurological examination (from unremarkable to variable degrees of cerebellar signs). Three patients presented an early-onset ataxia mimicking congenital ataxia with episodic syndromes. Early-onset non-progressive ataxia/hypotonia that might be isolated or might overwhelm episodic neurological manifestations represents a recently recognized mode of presentation of *CACNA1A* mutations,^{6,12,14,16,20} which has probably been underdiagnosed until recently.

Finally, our study also shows a large intrafamilial clinical heterogeneity with a classic presentation of well-known diseases related to *CACNA1A* (episodic ataxia type 2 and hemiplegic migraine) and a less typical presentation for which the diagnosis is often more difficult to make. Interviewing patients about their relatives can help establish the correct diagnosis and lead to appropriate care.

In conclusion, our results confirm that *CACNA1A* mutations are associated with a wide phenotypic spectrum including various forms of infantile or childhood onset episodic neurological events, neurological and cognitive abnormalities, and familial heterogeneity. The phenotypic spectrum associated with pathogenic mutations of *CACNA1A* includes non-epileptic paroxysmal events, paroxysmal epileptic manifestations, permanent neurological

symptoms, and cognitive disabilities. Identification of *CACNA1A* mutations must lead to close follow-up of psychomotor development and academic performance, so early management and care can be proposed. Patients with vermian atrophy are at a higher risk of cognitive dysfunction. We propose that a *CACNA1A*-associated phenotype may present as a neurodevelopmental disorder.

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SUPPORTING INFORMATION

The following additional material may be found online:

Appendix S1: Members of the Episodic Syndrome Consortium

Figure S1: Flow diagram illustrating the patients' inclusion

Figure S2: Features of the neurodevelopmental phenotype associated with *CACNA1A* mutations in childhood

Table SI: Diagnosis criteria of the episodic events

Table SII: Clinical and genetics characteristics of the patients

Table SIII: Clinical characteristics of the episodic events

Table SIV: Effects of the drugs on the episodic syndromes

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