Phenotypes of Spinocerebellar Ataxia Type 6 and Familial Hemiplegic Migraine Caused by a Unique CACNA1A Missense Mutation in Patients From a Large Family

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Background: Different mutations in the α1A-subunit of the brain P/Q-type calcium channel gene (CACNA1A) are responsible for familial hemiplegic migraine (FHM), episodic ataxia type 2, and spinocerebellar ataxia type 6 (SCA6). Missense and splice site mutations have been found in FHM and episodic ataxia type 2, respectively, whereas a CAG repeat in the CACNA1A gene was found expanded in patients with SCA6.

Objective: To identify the disease causing mutation in a large family of patients with phenotypes of hemiplegic migraine with or without cerebellar signs or permanent cerebellar ataxia without migraine inherited in a dominant manner.

Patients and Methods: We examined 15 patients from a large family identified through a systematic survey of hereditary ataxias being conducted in Portugal. Linkage analysis was performed with CACNA1A intragenic markers, and mutation analysis was performed by single strand conformational polymorphism analysis and sequencing.

Results: Genetic linkage analysis with CACNA1A intragenic markers showed positive LOD scores. The maximal LOD score was obtained with the polymorphic CAG repeat (Z_{max}=4.47, \theta=0). By single-strand conformational polymorphism analysis, a shift in exon 13 of the CACNA1A gene was detected in all patients. A G-to-A substitution was then identified, resulting in an arginine-to-glutamine change at codon 583 of this calcium channel α1A-subunit.

Conclusions: The disease-causing mutation in this family was identified, showing that a unique mutation in the CACNA1A gene causes several phenotypes, including those of SCA6 and FHM, thus suggesting that SCA6 and FHM are not only allelic diseases but are the same disorder with a large phenotypic variability.

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Familial Hemiclegic migraine (FHM) is a subtype of migraine with aura showing autosomal dominant inheritance. Episodes of FHM are characterized by some degree of hemiparesis occasionally associated with other symptoms, such as fever, drowsiness, confusion, or coma, which can be prolonged for days or weeks. Onset usually occurs during childhood or adolescence, although later onset has been reported. Permanent neurological signs of the disease are present in some patients, most often nystagmus and ataxia. Genetic heterogeneity of FHM has been established. A significant number of FHM families show genetic linkage to chromosome 19p13, including all those with cerebellar signs. Some FHM families without cerebellar signs have been assigned to chromosome 1p, whereas others are not linked to any known loci.

Episodic ataxia is a dominantly inherited paroxysmal cerebellar neurological disorder characterized by episodes of cerebellar ataxia, often accompanied by nausea, vertigo, and headache. Episodic ataxia type 1 (EA1) presents interictal myokymia during and between episodes due to mutations in a potassium voltage-gated channel gene, located on chromosome 12. Patients with episodic ataxia type 2 (EA2) show interictal nystagmus, and the disease gene maps to chromosome 19p. Migraine with or without aura may be present in some patients from EA2 families.

Spinocerebellar ataxias (SCAs) are progressive neurodegenerative disorders characterized by late-onset gait ataxia and dysthria. Seven dominantly inherited SCAs are caused by polyglutamine expansions: SCA1-2, Machado-Joseph disease, SCA6, SCA7, SCA17, and dentatorubropallidoluysian atrophy. The gene responsible for FHM, EA2, and SCA6 encodes an α1A-subunit of the brain P/Q-type calcium channel and is lo-
We studied a Portuguese family ascertained during a systematic, population-based survey of hereditary ataxias and spastic paraplegias, initiated in 1993 and covering half of the Portuguese population (5.6 million people). This family consisted of 17 patients with hemiplegic migraine and/or progressive cerebellar ataxia in 4 consecutive generations. Fifteen patients were clinically examined by one of us (P.C., J.B., or A.T.) (Table 1). Age at onset ranged from 3 to 23 years for migraine episodes (mean, 13.4 ± 7.2 years) and from 16 to 50 years for cerebellar ataxia (mean, 31.7 ± 11.5 years). The age at onset of migraine episodes varied from 8 to 71 years. Clinical manifestations consisted of 17 patients with hemiplegic migraine and/or progressive cerebellar ataxia, 15 of whom presented with phenotypes of either hemiplegic migraine or progressive cerebellar ataxia.24

To improve knowledge of the underlying mechanism involved in hemiplegic migraine and progressive cerebellar ataxia, we studied a large family in which patients presented with phenotypes of either hemiplegic migraine or progressive cerebellar ataxia, performed genetic linkage analysis with chromosome 19p13 markers, and performed mutation screening in the CACNA1A gene.

### METHODS

We studied a Portuguese family ascertained during a systematic, population-based survey of hereditary ataxias and spastic paraplegias, initiated in 1993 and covering half of the Portuguese population (5.6 million people). This family consisted of 17 patients with hemiplegic migraine and/or progressive cerebellar ataxia in 4 consecutive generations. Fifteen patients were clinically examined by one of us (P.C., J.B., or A.T.) (Table 1). Age at onset ranged from 3 to 23 years for migraine episodes (mean, 13.4 ± 7.2 years) and from 16 to 50 years for cerebellar ataxia (mean, 31.7 ± 11.3 years). The age at examination varied from 8 to 71 years. Clinical manifestations were pleomorphic, including episodes of altered consciousness precipitated by minor head trauma, focal neurological deficits precipitated or not by minor head trauma, and migraine without aura, besides progressive late-onset cerebellar ataxia in a few patients. One of these patients (III-3) was studied by brain magnetic resonance imaging, which showed atrophy of the cerebellum.

Peripheral blood samples were collected from patients and their relatives after written informed consent was obtained. Genomic DNA was obtained from peripheral blood leukocytes by standard techniques. Molecular analyses of genetic markers, exons, and intronic sequences of the CACNA1A gene were performed by polymerase chain reaction (PCR) amplification using the published primer sequences. The PCR was performed with 1 μM of each primer, 200 μM deoxynucleotides, 1 mM magnesium chloride, 10 mM Tris (pH 9.0), 50 mM potassium chloride, 1 μL of Taq polymerase, and 2% formamide in a final volume of 12.5 μL. The PCR products of markers were radioactively labeled and analyzed on 6% polyacrylamide gels. Allele sizes were determined by comparing migration relative to an M13 sequencing ladder.

Polyomorph markers, within a 4-centimorgan (M) interval containing the CACNA1A gene tel-D19S840-19S1150-(CAG)-D19S226-cen, according to the Fondation Jean Dausset Centre d’Etudes des Polymorphismes (Paris, France) database, were selected for linkage analysis. Markers D19S1150 and the polymorphic CAG repeat are intragenic. Analysis was performed with the LINKAGE software program version 5.22. The disease was considered autosomal dominant with incomplete penetrance (93%) and with a disease gene frequency of 0.0001. The PCR products of exons and intronic sequences of the CACNA1A gene were screened for molecular variants by single strand conformational polymorphism analysis and electrophoresis in ×0.5 Mutation Detection Enhancement gels (Bio-Whittaker Molecular Applications, Rockland, Me) at 4°C. Conformational changes were confirmed by sequencing with Thermosequenase cycle-sequencing kit (Amersham Pharmacia Biotech, Uppsala, Sweden). Restriction analysis of exon 13 was performed by PCR amplification, and products were digested with the BanII restriction enzyme (New England BioLabs, Beverly, Mass) according to manufacturer instructions.

### RESULTS

The size of the CAG repeat, in the 3’ end of the CACNA1A gene, responsible for SCA6 was determined in all 25 subjects available for this study. The repeat size in 12 patients, 9 at-risk individuals, and 4 spouses, ranged from...
In this study, we describe the first family to our knowledge in which patients presented phenotypes of hemiplegic migraine with or without cerebellar signs or permanent progressive cerebellar ataxia without migraine due to a unique missense mutation in the CACNA1A gene. The disease locus in this family showed strong linkage to intragenic markers in this gene. By mutation analysis, we identified an R583Q substitution in all available patients. This mutation had first been described in 2 affected members from a family with hemiplegic migraine and ataxia.<sup>3</sup> We described a large family with 17 patients who presented with high clinical variability due to this R583Q mutation.

The α<sub>1A</sub> subunit of the P/Q-type calcium channel gene is composed of 4 homologous domains (I-IV), each containing 6 putative transmembrane segments (S1-S6) and a pore-forming segment between S5 and S6.<sup>2</sup> The missense mutation identified in this family is located in the S4 transmembrane segment of protein domain II, which is thought to be the voltage sensor of the channel.

Mutation R583Q replaces a conserved, polar, positively charged arginine by a neutral glutamine, which can increase hydrophobicity and reduce polarity in this voltage sensor segment. This mutation causes a shift in the activation and inactivation voltage dependence of the channel to more negative potentials.<sup>29</sup> The hyperpolarization shift increases intracellular calcium levels by altering P/Q-type calcium channel activity at weak depolarizations in mutants with this substitution.<sup>29</sup> Channel recovery from inactivation in R583Q mutants is slower, which can lead to an accumulation of inactivated channels during rapid depolarizations.<sup>29</sup> Another FHM mutation due to an arginine-to-glutamate substitution also located in the S4 transmembrane segment, but of protein domain I at codon 192, also causes an excess of intracellular calcium due to altered gating properties.<sup>30</sup> The abnormal calcium influx, mostly during high neuronal activity, would explain the paroxysmal character of FHM and the precipitation of episodes by sensory or emotional stimuli.<sup>31</sup> Calcium overload causes excessive release of excitotoxic neurotransmitters such as glutamate, which can lead neurons to apoptotic death.

In this family, the mean age at onset for hemiplegic migraine symptoms was in the second decade and approximately 20 years earlier than that for the cerebellar signs. This onset of migraine symptoms is close to that reported in other clinical descriptions of FHM due to mutations in the CACNA1A gene.<sup>6</sup> The 2 patients previously described as having mutation R583Q began migraine episodes at 17 and 40 years, respectively, whereas cerebellar signs were first noticed in both patients when they were in their 60s.<sup>3</sup>

Emotional stress was the most frequent triggering factor of hemiplegic migraine in families with mutations in the CACNA1A gene as described in a previous study.<sup>6</sup> In the present family, patients with hemiplegic migraine did not refer to emotional stress as a triggering factor, whereas minor head trauma was referred to in approximately 4 patients (44%). However, this family is unique in which cerebellar progressive ataxia was also triggered by mild head trauma.

Expansion of a CAG repeat in the CACNA1A gene causes not only SCA6 but also EA2 phenotypes in patients from the same family.<sup>31,32</sup> On the other hand, in a family described by Yue et al.,<sup>24</sup> a point mutation in this gene originates severe progressive ataxia in some patients and episodic ataxia in others. Moreover, some families had members with either hemiplegic migraine accompanied by cerebellar signs or episodic ataxia with headache due to a point mutation in the CACNA1A gene.<sup>3,33,34</sup> In this family, we found patients who only had symptoms of progressive cerebellar ataxia, patients affected by hemiplegic migraine only, and patients with both hemiplegic migraine and symptoms of progressive cerebellar ataxia. Thus, the R583Q mutation causes phenotypes of SCA6 and FHM. These results, in addition to those referred to herein,<sup>31,32</sup>

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Table 2. Linkage Relationships Between the Disease Locus and Chromosome 19p13 Markers

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suggest that EA2, SCA6, and FHM are not only allelic diseases but are the same disorder with a large phenotypic variability. The presence of several different phenotypes strongly suggests the involvement of modifying polymorphisms in either this or other genes.

Mutations in the α1A subunit orthologous mouse gene are responsible for 2 phenotypes: the tottering (tg) and the leaker (tgla). The tgla mice phenotype presents severe progressive ataxia caused by a mutation in a splicing consensus sequence, which gives rise to CACNA1A aberrant transcripts. On the other hand, the tg mutant mice phenotype is caused by an amino acid substitution in the pore-forming region of mice α1A protein domain II. This mutant expresses a milder phenotype and shows less functional changes. The tg and tgla mutated channels exhibit a reduced calcium influx in Purkinje cells.

In conclusion, the mutation R583Q in the CACNA1A gene causes a large variety of clinical phenotypes, including hemiplegic migraine, permanent ataxia, and coma.
Mutations not only in the pore-forming segments but also in the voltage sensor transmembrane segments alter the gating properties of neuronal P/Q-type calcium channels, causing alterations in calcium influx through neurons.

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Author contributions: Study concept and design (Drs Silveira and Coutinho); acquisition of data (Ms Alonso, Drs Barros, Tuna, Silveira, and Coutinho, and Mr Coelho); analysis and interpretation of data (Ms Alonso, Drs Barros, Tuna, Sequeiros, Silveira, and Coutinho, and Mr Coelho); drafting of the manuscript (Ms Alonso, Drs Barros, Tuna, Sequeiros, Silveira, and Coutinho, and Mr Coelho); critical revision of the manuscript for important intellectual content (Ms Alonso and Drs Sequeiros, Silveira, and Coutinho); obtained funding (Dr Silveira); administrative, technical, and material support (Ms Alonso, Drs Barros and Tuna, and Mr Coelho); study supervision (Drs Sequeiros, Silveira, and Coutinho).

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