Progressive Myoclonus Epilepsy Caused by a Homozygous Splicing Variant of SLC7A6OS

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Exome sequencing was performed in 2 unrelated families with progressive myoclonus epilepsy. Affected individuals from both families shared a rare, homozygous c.191A > G variant affecting a splice site in SLC7A6OS. Analysis of cDNA from lymphoblastoid cells demonstrated partial splice site abolition and the creation of an abnormal isoform. Quantitative reverse transcriptase polymerase chain reaction and Western blot showed a marked reduction of protein expression. Haplotype analysis identified a ~0.85cM shared genomic region on chromosome 16q encompassing the c.191A > G variant, consistent with a distant ancestor common to both families. Our results suggest that biallelic loss-of-function variants in SLC7A6OS are a novel genetic cause of progressive myoclonus epilepsy.

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Progressive myoclonus epilepsies (PMEs) are a group of rare Mendelian disorders defined by the combination of action and reflex myoclonus, other types of epileptic seizures, and progressive neurocognitive impairment.1 PMEs usually begin in late childhood or adolescence. Disease progression is invariable, but prognosis and longevity vary considerably according to the etiology.

There are >20 known genetic causes of PME.2 They can be broadly divided into 2 clinical groups. The first group comprises those associated with significant cognitive decline, such as Lafora disease. The second group, having relatively preserved cognition, is largely exemplified by Unverricht–Lundborg disease (ULD), caused by biallelic mutations in CSTB.3

Despite recent advances, a substantial proportion of PME cases remain without a molecular diagnosis, with more genetic etiologies yet to be discovered.4,5 We report 2 families where multiple relatives are affected with a “ULD-like” PME phenotype and share the same rare homozygous splicing variant in SLC7A6OS (solute carrier family 7 member 6 opposite strand).

Patients and Methods
Ethical Approval and Exome Sequencing
Family 1 (Portuguese origin) was studied in a diagnostic setting and Family 2 (Turkish origin) in a research setting (Fig 1A). The institutional review board of Helsinki University Central Hospital approved the research study. Following exclusion of dodecamer repeat expansions in CSTB, paired-end exome sequencing (ES) was performed on 3 subjects (Family 1: IV-2; Family 2: V-4 and V-6; see Fig 1A). ES was performed on Nextseq 500 (150bp, Family 1) or HiSeq 2000 (75bp, Family 2; Illumina, San Diego, CA) following standard DNA enrichment protocols and analyzed using appropriate bioinformatic methods as previously described.6

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FIGURE 1: (A) Family pedigrees showing SLC7A6OS variant segregation and the type of genetic study performed in each subject. Consanguineous parental unions are indicated with 2 connecting lines. Consanguinity was not reported by the members of Family 1; however, the families were from the same rural area, and a common ancestor had been hypothesized. Subsequent inbreeding coefficient calculations supported the consanguineous unions as depicted by the dotted lines. (B) Migration on a Labchip GX of polymerase chain reaction products obtained by amplification of exons 1 to 3 of blood-derived SLC7A6OS cDNA. The expected fragment of 455bp was obtained for the 3 individuals tested (Patient III-2 in Family 1, his mother, and a control). For the patient and his mother, an additional 576bp amplicon was detected, representing, respectively, approximately 20% or 10% of the amount of SLC7A6OS cDNA, with the expected fragment decreasing in proportion. (C) Sanger sequencing of both the 455 and 576bp amplicons. The “abnormal” transcript corresponds to the retention of the complete intron 1 (+121bp). (D) In Western blot analysis, a decrease in the abundance of the SLC7A6OS protein was detected in the patient (III-2, Family 1) compared to controls. The analysis was performed in duplicates with 2 deposits each time. Mean estimations with a ChemiDoc imager were about 50% of the expression in controls. [Color figure can be viewed at www.annalsofneurology.org]
Expression Studies in Family 1

Blood-derived RNA samples were collected from the affected proband (IV-2) and his unaffected mother (III-2). cDNA was synthesized by standard techniques. Primers specific for “wild-type” SLC7A6OS transcript were used for real-time polymerase chain reaction (PCR): ex1/2F, TCCCAGGAGGAACCCGTCCA and ex2R, TCAGGTTCTCCCTCCTCGTG.

Proteins were extracted from lymphoblastoid cells. Forty micrograms of each sample was electrophoresed on NuPAGE 12% Bis-Tris Protein Gels (ThermoFisher Scientific, Waltham, MA) and transferred to Hybond-ECL Nitrocellulose membrane (GE Healthcare, Velizy-Villacoublay, France). The membrane was incubated overnight with primary antibodies: rabbit polyclonal anti-SLC7A6OS (ab122727; Abcam, Cambridge, UK) used at dilution of 1:250 or rabbit monoclonal antiactin (H&L (horseradish peroxidase; ab16284) in PBS with 3% milk and primary antibodies: donkey antirabbit IgG H&L (horseradish peroxidase; ab16284) in PBS with 3% milk. After washing, the membrane was incubated for 1 hour at room temperature with secondary antibodies: donkey antirabbit IgG H&L (horseradish peroxidase; ab16284) in PBS with 3% milk. Blots were revealed by ECL Prime kit (Pierce, Woburn, MA). The chemiluminescent signals of blots were quantified using a ChemilDoc imager (Bio-Rad, Marnes-la-Coquette, France).

Post Hoc Linkage and Haplotype Analyses

Nine family members from Family 1 and 6 from Family 2 were single nucleotide polymorphism (SNP) genotyped (see Fig 1A). Pairwise sample identity-by-descent (IBD) analyses were performed using KING, and inbreeding coefficients were estimated by FEstim. Autosomal recessive linkage analyses were performed using MERLIN. Genotyped SNP’s were manually interrogated to verify linkage breakpoints and determine haplotype lengths and sharing. Copy number variants (CNVs) were detected using PennCNV.

Brain Coexpression Exploratory Analysis

Gene coexpression analyses were performed using normalized developing brain expression data from BrainSpan (http://www.brainspan.org/). Genes were removed if they had expression values missing from >50% of samples or expression variation of <0.5 across all samples. A total of 15,957 genes, across 71 samples, remained in the filtered dataset. Using log2-transformed expression values, a matrix of weighted pairwise gene correlations was generated; weights were determined by $1/\sqrt{n}$, where $n$ is the number of samples contributed by the respective individual.

Results

Clinical Presentation of Patients

The clinical features of all 6 patients were similar and, in the early course of disease, resembled classic ULD (Table ). Electroencephalographic recordings showed relative preservation of background rhythms with generalized polyspike, polyspike–wave, and sometimes spike–wave discharges. Discharges were enhanced during hyperventilation and photic stimulation. Giant somatosensory evoked potentials were documented in patients from both families (Family 1: IV-2, Family 2: V-1). Brain magnetic resonance imaging at disease onset was normal (Family 1: IV-2, Family 2: V-6 and V-1), with serial studies in Family 2 demonstrating deterioration to marked cerebellar (V-6) and cerebral atrophy (V-1).

Exome Analysis Prioritizes SLC7A6OS Variant

Autosomal recessive modes of inheritance were hypothesized based on pedigree structure (see Fig 1A). Loss-of-function variants, including predicted splicing variants, were further prioritized. This approach led to the homozygous SLC7A6OS NM_032178.2: c.191A > G substitution being identified as the most likely pathogenic variant in both families (see Supplementary Table S1). Pedigree segregation supported the variant being disease-causing in the homozygous state (see Fig 1A).

Patient RNA and Protein Expression Studies

The SLC7A6OS c.191A > G affects the second last nucleotide of the first exon. Splicing prediction tools predicted abolition of the donor site (see Supplementary Table S1). Analysis of cDNA amplicons from exon 1 to exon 3 showed a single expected product of 455bp in controls (see Fig 1B). In Patient IV-2 and his mother III-2 (Family 1), the normal 455bp product was decreased in amount and there was an additional product of 576bp (see Fig 1B, C), resulting from the retention of the intron 1 (121bp), expected to lead to a frameshift.

Quantification of SLC7A6OS transcripts by quantitative reverse transcriptase polymerase chain reaction in Patient IV-2 revealed that expression of SLC7A6OS in blood was approximately 60% and in lymphoblastoid cells approximately 15% of that in controls. In the blood of the heterozygous mother (III-2), the SLC7A6OS expression was approximately 85% of controls. SLC7A6OS protein expression was also markedly reduced (see Fig 1D).

Post Hoc Linkage and Haplotype Analysis

Inbreeding estimates were consistent with consanguinity in both families (see Fig 1A). Under an autosomal recessive disease model, a single genome-wide significant linkage peak (logarithm of odds [LOD] > 3.0) was found for
both families on chromosome 16. No other regions with an LOD score of >0.5 were identified. The homozygous SLC7A6OS variant is located at chr16:68,344,639bp (86.6cM); both linkage regions include this base position (hg19; Fig 2A). No CNVs were detected in the linkage region for either family.

No close genetic relatedness between the 2 families was revealed by IBD analysis (third-degree relatives reliably excluded). However, to explore whether the variant may have arisen from a distant common ancestor, we compared the c.191A > G variant haplotype in both families. A total of 135 consecutive SNPs were shared by the 2 families at locus chr16q22.1 (chr16: 67,670,682–68,548,709; see Fig 2B). The shared haplotype is ~880kb (~0.85cM) in length, consistent with a founder effect.

### Brain Coexpression Exploratory Analysis

We explored brain coexpression patterns between SLC7A6OS and genes already known to cause an “ULD-like” PME phenotype (ie, CSTB, KCNC1, GOSR2,

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**TABLE. Clinical Presentation and Summary of Affected Individuals from Both Families with the c.191A > G Variant in SLC7A6OS**

<table>
<thead>
<tr>
<th>Patient ID</th>
<th>Family 1</th>
<th>Family 2</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>III-3</td>
<td>III-4</td>
</tr>
<tr>
<td>Sex</td>
<td>F</td>
<td>M</td>
</tr>
<tr>
<td>Development</td>
<td>Normal</td>
<td>Normal</td>
</tr>
<tr>
<td>Age at onset</td>
<td>21 yr</td>
<td>17 yr</td>
</tr>
<tr>
<td>First symptom</td>
<td>TCS</td>
<td>TCS</td>
</tr>
<tr>
<td>Myoclonus</td>
<td>+++</td>
<td>+++</td>
</tr>
<tr>
<td>TCS</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Cerebellar signs</td>
<td>Ataxia</td>
<td>Ataxia</td>
</tr>
<tr>
<td>Cognitive decline; dysfunction</td>
<td>No</td>
<td>No</td>
</tr>
<tr>
<td>Outcome (age)</td>
<td>Wheelchair (22 yr); living (41 yr)</td>
<td>Wheelchair (21 yr); living (38 yr)</td>
</tr>
<tr>
<td>Antiseizure drugs</td>
<td>CNZ, LRZ, LEV, VPA</td>
<td>CNZ, DZP, LEV, VPA</td>
</tr>
<tr>
<td>Other diagnoses</td>
<td>Depression (mild)</td>
<td>Depression (mild); addiction (drugs and alcohol)</td>
</tr>
</tbody>
</table>

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*Walking acquired after the age of 2 years.

bFebrile seizure at 1 year.

cPossible afebrile seizure at the age of 1.5 years and poor balance noted at 9 years.

dSymptom aggravation.

+ = mild; ++ = moderate; +++ = severe; − = not observed; CLB = clobazam; CNZ = clonazepam; DZP = diazepam; F = female; LEV = levetiracetam; LRZ = lorazepam; M = male; PHT = phenytoin; PRIM = primidone; TCS = tonic-clonic seizures; TPM = topiramate; VPA = valproate; ZNS = zonisamide.
SCARB2, ASAH1, CERS1). The analysis revealed that SLC7A6OS and SCARB2 are in the top 5% of the most highly coexpressed genome-wide genes in the developing brain; SLC7A6OS and GOSR2 are in the top 10%. With ASAH1, the 4 genes form a set of significantly positively correlated genes that all negatively correlate with CERS1 ($p < 0.01$, Monte Carlo sampling).

**Discussion**

We describe 2 unrelated families with PME and an identical homozygous variant affecting the splicing of SLC7A6OS. The similar phenotypic presentation of the 6 patients, combined with a likely founder effect origin of the variant, strongly suggests that it is the underlying cause of PME for these patients. Whereas the clinical and
electrophysiological presentation was similar to ULD.\textsuperscript{3,12} Onset age was slightly later (11–21, mean = 14 years) and progression with loss of ambulation and cognitive decline was faster than typical for ULD.

The c.191A > G variant was predicted to disrupt an SLC7A6OS donor splice site and patient CDNA, and protein studies confirmed this. We observed the production of an abnormal isoform with the retention of intron 1. Expression of SLC7A6OS in blood was approximately 60% and in lymphoblastoid cells approximately 15% of that in controls. Expression of the normal SLC7A6OS isoform was markedly reduced, which was confirmed by a decrease of ~50% of the SLC7A6OS protein product.

The exact function of SLC7A6OS remains unknown, although there is some evidence to support it playing a role in RNA polymerase II nuclear import.\textsuperscript{13} In the zebrafish, slc7a6os is highly expressed in the central nervous system (CNS) during development.\textsuperscript{14} Furthermore, functional knockdown of slc7a6os by splice-blocking morpholinos showed morphological defects in the CNS and reduced mobility. Interestingly, the morpholino used by Benini and colleagues\textsuperscript{14} targets the slc7a6os exon1–intron1 boundary and is expected to cause the retention of intron 1 in the mature mRNA, thus mimicking the effect of the c.191A > G variant in our PME patients.

To further support the role of SLC7A6OS in PME, we compared its coexpression patterns in the developing human brain with established “ULD-like” disease genes. Our analysis identified SCARB2 as the established “ULD-like” gene, with the most highly correlated brain coexpression pattern with SLC7A6OS, and that together with GOSR2, ASAH1, and CERS1 they form a cluster of significantly coexpressed genes. With a putative role for SLC7A6OS in nuclear import, all 5 genes are broadly involved in intracellular trafficking, with correlated coexpression signatures supporting a potential shared pathway in the brain specifically.

In conclusion, these data suggest that biallelic variants in SLC7A6OS, displaying a founder effect, are a rare, new cause of PME, with a phenotype similar to ULD.

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**Author Contributions**

Conception and design of the study: G.L, A-E.L, S.F.B., M.B. Acquisition and analysis of data: all authors. Drafting a significant portion of the manuscript or figures: L.M., K.L.O., A.L., S.F.B., A-E.L., G.L.

**Potential Conflicts of Interest**

Nothing to report.

**References**