Loss-of-function variants in the KCNQ5 gene are implicated in genetic generalized epilepsies

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Summary

Background De novo missense variants in KCNQ5, encoding the voltage-gated K+ channel KV7.5, have been described to cause developmental and epileptic encephalopathy (DEE) or intellectual disability (ID). We set out to identify disease-related KCNQ5 variants in genetic generalized epilepsy (GGE) and their underlying mechanisms.

Methods 1292 families with GGE were studied by next-generation sequencing. Whole-cell patch-clamp recordings, biotinylation and phospholipid overlay assays were performed in mammalian cells combined with homology modelling.

Findings We identified three deleterious heterozygous missense variants, one truncation and one splice site alteration in five independent families with GGE with predominant absence seizures; two variants were also associated with mild to moderate ID. All missense variants displayed a strongly decreased current density indicating a loss-of-function (LOF). When mutant channels were co-expressed with wild-type (WT) KV7.5 or KV7.5 and KV7.3 channels, three variants also revealed a significant dominant-negative effect on WT channels. Other gating parameters were unchanged. Biotinylation assays indicated a normal surface expression of the variants. The R359C variant altered PI (4,5)P2-interaction.

Interpretation Our study identified deleterious KCNQ5 variants in GGE, partially combined with mild to moderate ID. The disease mechanism is a LOF partially with dominant-negative effects through functional deficits. LOF of KV7.5 channels will reduce the M-current, likely resulting in increased excitability of KV7.5-expressing neurons. Further studies on network level are necessary to understand which circuits are affected and how this induces generalized seizures.

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Introduction

KCNQ5 is one of five members of the highly conserved KCNQ gene family, which encodes the α-subunits of the M-type, voltage-gated delayed rectifier potassium channels K_7.1-7.5. Four subunits can form a functional channel as either homo- or heterotetramers. Heteromeric channels containing K_7.3 subunits together with K_7.5 or K_7.2 have been shown to yield larger K⁺ currents than each of the subunits alone. The KCNQ1 gene is mainly expressed in cardiac muscle and the cochlea, KCNQ2 and -3 are mainly expressed throughout the central nervous system (CNS) and peripheral nervous system (PNS). KCNQ4 is expressed in sensory outer hair cells and KCNQ5 is expressed in the brain, skeletal muscle and blood vessels. In addition to heteromerization, native Kv7 currents, particularly those of K_7.1, can be influenced by assembly with KCNE subunits and variants in either one of them can cause cardiac arrhythmia or deafness. Due to its very restricted expression, KCNQ4 is a key contributor to the auditory system and its dysfunction has been associated with non-syndromic dominant deafness.

The other three family members, KCNQ2, -3 and -5, are important regulators of the neuronal M-current within the nervous system that very effectively controls neuronal firing. In particular, dominant negative K_7.5 channel expression has been found to decrease the medium and slow afterhyperpolarization currents in the CA3 region of the hippocampus in a mouse model. Moreover, this model unravelled the role of KCNQ5 in attenuating synaptic inhibition and modifying hippocampal network synchronization. Upon application of muscarinic receptor agonists, such as acetylcholine, a signalling cascade is triggered that causes depletion of phosphatidylinositol-4,5-bisphosphate (PI (4,5)P₂) from the membrane, which forces Kv7 channels to close reducing the M-current and resulting in increased neuronal firing.

Loss-of-function (LOF) variants in KCNQ2 and KCNQ3 were first identified as the cause of epileptic seizures in patients with benign familial neonatal epilepsy (BFNE) and later in developmental and epileptic encephalopathies (DEE). More recently, de novo heterozygous missense and truncating variants in KCNQ5 in individuals with intellectual disability (ID)
alone or with DEE have been described.\textsuperscript{16,17,18} Additionally, an individual presenting with absence seizures in adolescence, migraine and mild ID has recently been identified with an intragenic duplication of KCNQ5 most likely causing haploinsufficiency by skipping exons 2-11 and resulting in a premature stop codon.\textsuperscript{19}

Here, we analysed different cohorts of GGE patients to identify causative variants in KCNQ5, studied the phenotypes of affected individuals and co-segregation of the detected variants. Electrophysiological and biochemical characterization of variants contributing to the disease was performed by expression of mutant K\textsubscript{\textit{v}}7.5 subunits in Chinese hamster ovary (CHO) cells and using whole cell patch-clamping, biotinylation and phospholipid overlay assays.

**Subjects/materials and methods**

**Study participants**

This study was approved by the local institutional review boards of the participating centres. The patients or their relatives gave their written informed consent. In total, 10 individuals from 5 families were ascertained from three different cohorts. Individuals 1, 2, 3, 4, 6 and 7 were ascertained from the EuroEPINOMICS-CoGIE study, while individual 5 was ascertained from the Epi25 study (Tuebingen subcohort), individuals 8 and 9 were analysed using a gene panel in a French cohort, and individual 10 was ascertained from the EPGP/Epi4K study. Two additional independent individuals from the latter cohort carried benign KCNQ5 variants and are not considered further in our study (compare first paragraph of the results section). Medical and family histories, neurological examination, brain imaging and EEG findings were analysed. Seizure types were classified according to the latest International League Against Epilepsy classification.\textsuperscript{20} Blood samples were taken from available family members and DNA was extracted by standard procedures.

**Genetic analysis**

DNA from individuals from all cohorts were whole exome sequenced, whereas individuals 8 and 9 were analysed in an epilepsy gene panel. Validation of discovered KCNQ5 variants was performed via Sanger sequencing with the exception of family 3 due to insufficient amount of DNA. Splice site variants were further investigated by RNA extraction from patient blood (PAXgene Blood RNA System, BD), and subsequent cDNA (NM\textsubscript{_019842.3}) were purchased from GeneScript (Netherlands). Site-directed mutagenesis was performed using PCR with Pu\textsubscript{i} polymerase (Promega, Germany). The inserts were sequenced to confirm the introduction of the point mutations and to exclude additional alterations.

**Variant Interpretation**

KCNQ5 variants were considered putatively disease-relevant if 3/4 of the following criteria were met (i) likely functional effect (protein-truncating variant, inframe-deletion or a missense variant with at least 4 pathogenic predictions), (ii) minor allele frequency (MAF) of 0 in the European populations of the 1000 genomes (http://www.1000genomes.org), the Exome Variant Server (EVS; http://evs.gs.washington.edu) and in the Genome Aggregation Database (gnomAD; http://gnomad.broadinstitute.org), (iii) confirmed in all affected family members, and (iv) the variant demonstrated an abnormal effect in electrophysiological recordings or analysis of splicing. The variants were interpreted according to the American College of Medical Genetics and Genomics standards and guidelines for the interpretation of sequence variants.\textsuperscript{21}

**Testing for variant enrichment**

To estimate the burden of qualifying variants in KCNQ5 in GGE cases vs. controls, we extended the rare-variant association analysis performed by Epi4K/EPGP\textsuperscript{22} using additional cohorts of unrelated GGE patients and matched controls, all of European ancestry. A combined non-overlapping set of 4,418 GGE cases and 7,727 controls from these previously published cohorts was investigated using the Cochran-Mantel-Haenszel exact test as follows: 1) 640 cases and 3,877 controls from the Epi4K/EPGP study,\textsuperscript{22} 2) 874 cases and 2,177 matched controls from EuroEPINOMICS-CoGIE, EpiPGX & CENet consortia studies,\textsuperscript{23-24} 3) 2,904 cases and 1,763 matched controls from the Epi25 Collaborative study.\textsuperscript{25} Qualifying variants were defined consistently across cohorts using the criteria previously defined by Epi4K/EPGP.\textsuperscript{22}

**Functional analysis**

**Mutagenesis.** pcDNA3.1-P2A-eGFP and pcDNA3.1-P2A-taRFP vectors containing the human K\textsubscript{\textit{v}}7.5 subunit cDNA (NM\textsubscript{_019842.3}) were purchased from GeneScript (Netherlands). Site-directed mutagenesis was performed using PCR with Pu\textsubscript{i} polymerase (Promega, Germany). The inserts were sequenced to confirm the introduction of the point mutations and to exclude additional alterations. A pcDNA3-PIP3K\textsubscript{\textit{V}} (NM\textsubscript{_001146687.2}) plasmid was kindly gifted by Alvaro Villarroel (Instituto Biofisika, University of Basque Country, Leioa, Spain). pDrVSP-IRES2-EGFP was a kind gift from Yasushi Okamura (Addgene plasmid # 80333; http://n2t.net/addgene:80333; RRID: Addgene\_80333).

**Transfection and expression in CHO cells.** CHO-K\textsubscript{1} cells were cultured at 37 °C in a 5% CO\textsubscript{2} humidified atmosphere and grown in Ham’s F12 containing 2mM glutamine (Gibco), 10% (v/v) fetal calf serum (PAN Biotech; Tuebingen cohort) or a 1:1 Dulbecco’s modified Eagle medium (DMEM)/Ham’sF12 mix including 10% (v/v) fetal calf serum and 1% Penicillin-Streptomycin.
(Gibco; + antibiotics cohort) in 3.5 cm plastic dishes. Transfection was performed 24–48 h prior to electrophysiological recordings with Lipofectamine 3000 (Invitrogen) following the manufacturer’s protocol using 2-2.5 µg DNA and an additional 0.25 µg of eGFP DNA for the truncation variant (pA301Gfs*64) lacking the fluorescent marker due to the early stop codon. For co-transfection with WT subunits the same protocol was applied using 2–2.5 µg of DNA in total in a molar ratio of 1:1. For the WT controls only 1 µg of DNA (P2A-tagRFP construct) was used. For co-expression of Kv7.3, a CHO-K1 line stably expressing Kv7.3 channels was DrVSP or PIP5K plasmid.

KCNQ5 or augmentation, cells were co-transfected with 2 µg of either the WT, mutant cDNAs or water (mock) using the following bath solution contained (in mM): 138 NaCl, 2 CaCl2, 5.4 KCl, 1 MgCl2, 10 glucose and 10 (4-[2-(Hydroxyethyl)-1-piperazineethanesulfonic acid (HEPES) (pH 7.4 adjusted with NaOH). Borosilicate glass pipettes had a tip resistance of 1.5–3.5 MΩ and were filled with pipette solution containing (in mM): 140 KCl, 2 MgCl2, 10 EGTA, 10 HEPES, 5 K2ATP (pH 7.4 adjusted with KOH).26 All recordings were performed at room temperature of 21–23 °C (RT).

Cells were visualized using an inverted microscope (Axio-Vert.A1, Zeiss or Nikon Eclipse). In case of single plasmid transfections only green (eGFP) or red (tagRFP for the WT in heteromeric expression experiments) fluorescent cells were selected for electrophysiological recordings 24–48 h after transfection, whereas in co-transfection recordings, cells were selected that showed an approximately equal amount of both, red and green fluorescence.

**Electrophysiology.** Standard whole-cell patch clamp recordings were performed using an Axopatch 200B or Multiclamp 700B amplifier, a Digidata 1320A, 1440A or 1550B digitizer (Axon Instruments), and pCLAMP 8, 10.4 or 11.1 data acquisition software (Molecular Devices). Leakage and capacitive currents were automatically subtracted using a pre-pulse protocol (-P/4). Cells were held at -80 mV in whole-cell configuration for 2 min prior to recording and series resistance was compensated (at approx. 85%) and monitored regularly. Currents were filtered at 1 kHz and digitized at 5 kHz. The bath solution contained (in mM): 138 NaCl, 2 CaCl2, 5.4 KCl, 1 MgCl2, 10 glucose and 10 (4-[2-(Hydroxyethyl)-1-piperazineethanesulfonic acid (HEPES) (pH 7.4 adjusted with NaOH). Borosilicate glass pipettes had a tip resistance of 1.5–3.5 MΩ and were filled with pipette solution containing (in mM): 140 KCl, 2 MgCl2, 10 EGTA, 10 HEPES, 5 K2ATP (pH 7.4 adjusted with KOH).26 All recordings were performed at room temperature of 21–23 °C (RT).

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**Patch clamp protocols and data analysis.** K+ currents were induced by depolarizing the membrane from a holding potential of -80 mV to +60 mV in 10 mV steps for 2 s. Subsequently, a shorter hyperpolarizing pulse was elicited to −120 mV for 0.5 s to obtain tail currents. Current amplitudes were calculated from the mean steady-state current for the last 0.5 s of the first step depolarization. Current densities (pA/pF) were obtained by normalizing the current amplitudes to the cell membrane capacitance. The activation curve was determined by plotting the normalized tail (Itail) current against the step potential (Vj). A Boltzmann function, ITail = 1 / (1 + exp[(V0.5 − Vj)/k]), where V0.5 is the voltage of half-maximal activation and k is the slope factor, was fit to the data points. In experiments using voltage-sensing phosphatase, a step from −80 mV to 0 mV was applied for 2 s followed by a step to +100 mV for 0.2–2.0 s to activate the phosphatase before decreasing membrane voltage back to 0 mV for 25 s. Data from recordings using voltage-sensing phosphatase were analysed by normalizing the negative peak to the mean current before the step to +100 mV for each recording and results were fit with a sigmoidal function. Furthermore, the recovery of current over time was analysed by normalizing the current at each second after the step to the mean current prior to the step. Clampfit software of pClamp10.7 (Axon Instruments), Microsoft Excel (Microsoft Corporation, Redmond, WA, USA) or GraphPad software (GraphPad Prism 8, San Diego, CA, USA) were used for data and statistical analysis. All data is shown as mean ± SEM. All data were tested for normal distribution. One-way ANOVA with Dunnett’s post hoc test or Student’s unpaired t-test were used to evaluate statistical significance of normally distributed data. If the data was not normally distributed, a Kruskal-Wallis test was performed followed by a Benjamiini, Krieger, and Yekutieli test. For all statistical tests p < 0.05 was considered significant. Scatter plots show single cell values, median and interquartile range.

**Western blot analysis.** CHO cells were lysed 24 h after transfection with either wildtype KCNQ5, one of the mutant cDNAs or water (mock) using the following buffer (in mM): 20 Tris (pH 7.5), 150 NaCl, 1 EDTA, 1 EGTA, 2.5 Napyrophosphate, 1 β-glycerophosphate, 1 sodium-orthovanadate, 10 DTT, 1% Triton and 0.05% protease inhibitor cocktail solution (Roche). Total protein concentration was measured via Bradford assay. 8% polyacrylamid gels were loaded with 20 µg of total protein per lane to separate these by sodium dodecyl sulfate-polyacrylamid gel electrophoresis (SDS PAGE). After transferring the proteins onto a nitrocellulose membrane (Whatman) via electrophoresis at 4 °C in Towbin buffer (25 mM Tris, 192 mM glycine, pH 8.3, 10% (v/v) methanol), the blots were blocked in 5% non-fat dry milk powder in phosphate-buffered saline with 1% Tween (PBST) for 1 h at RT. Subsequently, membranes were probed with a polyclonal rabbit primary antibody against Kv7.5 (ABN1372, Millipore) at 1:7,500 and a monoclonal mouse primary antibody against actin.
Analysis was performed using ImageJ 1.52r.27

Miniscence (ECL; Amersham, Cytiva). Quantitative
PBST, detection was performed via enhanced chemiluminescence (ECL; Amersham, Cytiva). Quantitative analysis was performed using ImageJ 1.52r.27

Biotinyllation assay. For isolation of membrane proteins, the Pierce Cell Surface Protein Biotinylation and Isolation Kit (ThermoFisher) was used according to the manufacturer’s protocol. In brief, cells were cultured for 48 h and transfected with either the WT or one of the mutant plasmids. Cells were biotinylated, lysed and isolated by binding to probed agarose beads 24 h after transfection. After elution, the proteins were prepared for Western blot analysis. Western blots were performed and analysed as described above. A monoclonal mouse anti-actin primary antibody (1:30,000, A5441, Sigma-Aldrich) at 1:30,000 overnight at 4 °C.

Protein-phospholipid overlay assay. PIP strips (Molecular Probes) were used as described in the manufacturer’s protocol. Concisely, cells were cultured and transfected as described above and lysed 24 h after transfection. PIP strips were blocked in 3% fatty-acid free bovine serum albumin (BSA) in Tris-buffered saline with 1% Tween (TBST) for 1 h at RT. Next, membranes were incubated at 4 °C overnight in TBST+3% BSA and a final protein concentration of 1 µg/ml. Membranes were washed and treated, developed and analysed the same way as Western blots described above.

Molecular modelling. Molecular modelling was performed using YASARA Structure, ver. 21.12.19 and OriginPro 2022 for data analysis. The Kv7.5 homology model was calculated based on the closely related Kv7.4 structure in complex with PIP(4,5)P2 7VNP.pdb.28 Five alternative alignments of Kv7.4 vs. 7.5 were generated with a maximum allowed (PSI)-BLAST E-value to consider a template (EValue Max) of 0.1. PSI-BLAST was applied to create a target sequence profile and feeding it to the PSI-Pred secondary structure prediction algorithm.29 Five Kv7.5 homology models were calculated based on the alignments. The homology modelling parameters were: modelling speed (slow = best); Slow; maximum oligomerization state (OligoState): 4 (tetrameric); maximum number of conformations tried per loop (LoopSamples): 50; maximum number of residues added to the termini (TermExtension): 10. Using YASARA Structure a consensus Kv7.5 homology model covering Kv7.5 residues 92-586 was generated. This model was employed for further Kv7.5 control simulation. To generate two slightly different start conformations, the model was energy minimized a second time and the resultant slightly different model was used for control simulation 2. To generate a Kv7.5 structure complexed to PIP(4,5)P2, the PIP(4,5)P2 molecules present in the template Kv7.4 structure 7VNP.pdb were positioned in virtually identical position into the two Kv7.5 consensus homology models. Subsequent energy minimization was applied to generate the Kv7.5-PIP(4,5)P2 models used for further simulations in presence of PIP2. The energy minimization procedure included an initial local steepest decent minimization without electrostatics to remove bumps followed by a simulated annealing minimization to reduce the energy of the Kv7.5-PIP2 complex. Kv7.5-R359C and Kv7.5-R359C-PIP2 complexes were generated by swapping arginine 359 to cysteine in the individual models followed by energy minimizations. Molecular dynamics simulation of the WT or mutant Kv7.5 or Kv7.5-PIP2 membrane proteins were run using force field AMBER14. Simulation temperature was set to 298 K and pressure was set to 1 bar at an electrostatics cutoff=8. The ion concentration in the solution was 0.9% NaCl (mass fraction) at pH 7.4. The channel models were embedded in phosphatidyl-ethanolamine (PEA) membranes in a square shaped simulation box that was 20 Å larger than the protein (periodic boundary). Membrane insertion of channels was performed using YASARA Structure standard procedure that scans the protein for putative transmembrane segments and positions the protein accordingly into the membrane. Membrane density of “1” is achieved by firstly insertion of a xy-directional-shrunken model into a central hole pinched in the membrane and subsequent extension plus MD simulation of the protein-membrane to normal size. MD simulation on the resultant system was initiated by an equilibration period of 250 picoseconds. During this initial equilibration phase, the membrane is artificially stabilized to allow for realistic repacking and cover the solute, while solvent molecules H2O, Na+ and Cl− are kept outside of the membrane region. The following MD simulation was computed as all-atoms-mobile simulation. All eight simulations were run for 100 nsec. The simulations were run on a 32-core AMD Ryzen Threadripper 2990WX computer equipped with 4 GeForce 2080 Ti graphic cards installed. As the modelling systems were relatively extensive (about 350.000 atoms) the individual simulation took about 6-8 weeks each. Root mean square deviation (RMSD), RMSF and Ca-cross-distances were calculated with provided YASARA macros md_analyze, mcr or manually. Where relevant, significance of mean differences for simulation data was tested by paired
Student’s t-test conditions within simulation 1 or simulation 2 indicated by * for $p < 0.05$.

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The funders had no role in study design, data collection, analyses, and interpretation of the data or in the writing of this publication.

Results

Genetic Screening of GGE Cohorts
In total, we identified 10 individuals within 5 families with a GGE phenotype carrying a disease-associated variant (see criteria in methods) in KCNQ5. No other variants in genes that have been associated with epilepsy before were detected. R359C and Q735R were first detected, each in 1 of 238 independent GGE families from the EuroEPINOMICS-CoGIE cohort.22 Both variants co-segregated in 2-4 affected family members and were found each also in one asymptomatic carrier. This prompted us to search for further variants in three other cohorts. L692V was found in 1 individual of 339 GGE families in the Epi25 subcohort from Tuebingen, E265_T306del in 2 related individuals in an epilepsy gene panel that was performed in 75 individuals in France, and A301Gfs*64 in 1 individual of 640 GGE families from the Epi4K/EPGP cohort.23 Two additional variants from the Epi4K/EPGP cohort, F165I and L926S, were found one time each in the gnomAD or TOPMed databases and did not show any alterations in electrophysiological recordings as compared to the WT (see Table 2). Hence, the individuals carrying these benign variants will not be considered further on. The pedigrees of families 1-5 are shown in Figure 1A.

Statistical analysis was performed to validate KCNQ5 as a potentially predisposing gene for GGEs. The odds of the occurrence of qualifying variants (as previously defined by the Epi4K/EPGP22) in KCNQ5 in GGE cases vs. controls was calculated over three large previously published cohorts (partially overlapping with the discovery cohorts mentioned above), each matched with independent controls. In total, we identified 8 qualifying variants in cases vs. 3 in controls, with a stratified odds ratio of 10.6 (confidence interval: 2.4-64.8; $p = 0.00044$). The odds were homogeneous across cohorts (Breslow-Day test; $p = 0.9$). This suggests that KCNQ5 variants are enriched in GGE cases, considering this hypothesis-driven approach.

Clinical descriptions

Family 1 (R359C). The family of German descent includes four clinically affected individuals with variable manifestations. Whereas the mother and the son suffer from absence seizures only, one daughter exhibits myoclonic seizures and generalized tonic-clonic seizures (GTCS) in addition to absences. The other affected daughter (individual 3) presents with GTCS alone. Age at epilepsy-onset varied between 3 and 7 years. All three affected children have ID. However, while individuals 2 and 3 already showed a developmental delay or ID before the onset of epilepsy, ID in individual 1 started after the onset of epileptic seizures. The mother has a normal intellect. One non-affected sibling carries the variant too, indicating reduced penetrance. Seizure control differs between the individuals of family 1 though lamotrigine seemed to have a positive effect in all affected family members. Relevant comorbidities have only been observed in the most severely affected individual (individual 1), who suffers from early onset arterial hypertension, obesity and mild generalized cerebellar ataxia.

Routine-EEG in individual 1 and 2 revealed generalized irregular 3/s (poly-)spikewave complexes, photoparoxysmal reaction and eyelid myoclonia under photostimulation. Generalized spike waves have also been present in individual 3. EEG recordings of individual 4 were not available. Cerebral imaging with MRI or CT-scan has not been performed in any of the family members. Clinical features of all our individuals are summarized in Table 1.

Family 2 (L692V). Individual 5 is the only affected family member in family 2 and is of Chechen origin. She started to have absence seizures at the age of seven years, and later had GTCS. She is not seizure-free and never took anti-seizure medication (ASM) regularly, so we cannot comment on ASM response. Development was normal. She also suffered from migraine with aura. Physical examination was normal. In the EEG she presented generalized spike-waves and MRI was described as normal. Paternal genotypes could not be established as the trace to the parents has been lost during the Chechen war.

Family 3 (Q735R). In this consanguineous Turkish family (parents are first-degree cousins), two affected brothers were diagnosed with juvenile myoclonic epilepsy (JME) with a typical age of onset (14 or 16 years) of myoclonic seizures, and later GTCS. One of the brothers (individual 7) also presented with absence seizures. The variant was inherited from the unaffected mother. For individual 6 generalized epileptiform discharges were recorded on EEG. Individual 7 did not undergo EEG or cerebral imaging, while a cerebral CT scan of individual 6 was normal. Diagnosis of individual 7 was based on seizure description and positive family history. Development and neurological examinations were normal in both individuals. Individual 6 is pharmaco-resistant, whereas individual 7 is not (more definitive information on ASM regimen was not available).
Family 4 (E265-T306del). Both affected individuals of the French family started to have absence seizures at the age of one and a half years (18 and 19 months, respectively). The mother (individual 9) also suffered from GTCS. They share a similar phenotype since both present with mild ID and similar age of onset (see Table 1). Individual 8 still has rare absences, individual 9 is seizure-free under medication. Neurological examination and MRI scans were normal in both. The EEG of individual 8 showed generalized spike-waves, the EEG of individual 9 was normal.

Family 5 (A301Gfs*64). Individual 10, Australian born and of European descent, was diagnosed with JME (myoclonic seizures, absences, GTCS) at the age of 14 years. While neurological examination, cognitive status and MRI were normal, the EEG showed polyspike-slow-wave patterns and photoparoxysmal responses. She is seizure-free under medication (see Table 1). Interestingly, her twin sister, had a single GTCS induced by flashing lights in a discotheque at the age of 19 years. The twin sister refused genetic testing.

Functional characterization of KCNQ5 variants. The KCNQ5 c.918+5G>A (E265_T306del) variant in individuals 8 and 9 affects a donor splice-site. Splicing prediction tools predicted abolition of the donor site (SpliceAI score 0-98). To examine the consequences of this variant, RNA was extracted from blood, and analysis of cDNA amplicons with primers located in exon 2 and at the junction of exon 6 and 7 was performed and revealed the expected 583 bp band for the wild type (WT), and an additional smaller band of 460 bp, corresponding to the complete skipping of exon 5. This was confirmed by sequencing of the PCR products. To this end, the variant was assumed a LOF due to the large deletion within the critical pore region of the channel (S5 segment; see Figure 1B).

To functionally characterize the remaining variants, CHO cells were transfected either with the mutant KCNQ5 cDNA and compared to those transfected with WT KCNQ5 cDNA (2-2.5 µg), or with a 1:1 mix of mutant and WT cDNA (1 µg each) and compared to the same amount of WT cDNA alone (1 µg) to test for a dominant-negative effect of mutant on WT channels.
| Family | Descendant | Epilepsy Syndrome | Variant | Functional consequence | Consanguinity | Sex, age at investigation | Age of onset (at onset) | Seizure type (at onset) | Seizure outcome (2020) | Development before seizure onset | Development | Previous or active medication | Comorbidities | Neurological examination | EEG | MRI/CT |
|--------|------------|-------------------|---------|------------------------|--------------|------------------------|-----------------------|------------------------|------------------------|-------------------------|--------------|-----------------------------|--------------|-----------------------------|-----|--------|
| 1      | Individual 1 | German            | CAE→JME | 1075C>T               | No           | Female 19y              | 3y                    | Absence, Myoclonus     | Seizure free with VPA+LTG+ETX | Mild developmental delay starting between 3,5y and 5 y | Moderate ID | Received VPA 25 mg/kg, outcome unknown | Arterial hypertension, Obesity | Mild ataxia | GSW, PPR (12Hz) gen. irreg. 3/s poly SW for 15sec, eyelid myoclonia under phs | n/a | n/a |
| 2      | Individual 2 | German            | CAE     | 1075C>T               | No           | Male 6y                 | 5y                    | Absence               | Seizure free with LTG | Normal | VPA+LTG | None | Normal | GSW |
| 3      | Individual 3 | German            | IGE-GTCS | 1075C>T              | No           | Female 14y              | 7y                    | Absence, GTCS         | Seizure free with LTG | Normal | VPA+LTG | None | Normal | GSW |
| 4      | Individual 4 | German            | CAE     | 2074C>G               | No           | Female 43y              | 7y                    | Absence, GTCS         | Seizure free with LTG | Normal | VPA+LTG | None | Normal | GSW |
| 5      | Individual 5 | Chechen           | CAE→JME-CAE-IGE-GTCS | 2074C>G     | Male 27y                | Female 37y             | 7y                    | Absence               | Seizure free with LTG | Normal | VPA+LTG | None | Normal | GSW |
| 6      | Individual 6 | Turkish           | CAE     | 2204A>G               | No           | Male                    | 14y                   | Myoclonus, GTCS       | Not seizure free due to incompliance (VPA+LEV) | Mild ID | Mild ID | VPA+LEV | Mild ID | GSW |
| 7      | Individual 7 | Turkish           | JME     | 918+5G->A             | No           | Male                    | 16y                   | Myoclonus, GTCS       | 2-3 myoclonic seizures/week with VPA+TPM | Mild ID | Mild ID | LEV | Mild ID | GSW |
| 8      | Individual 8 | French            | JME     | 918+5G->A             | No           | Male                    | 18m                   | Abnormal              | Seizure free with LEV | Normal | VPA+LEV | None | Mild ID | GSW |
| 9      | Individual 9 | French            | CAE     | 918+5G->A             | No           | Male                    | 19m                   | Abnormal              | Seizure free with LEV | Normal | VPA+LEV | None | Mild ID | GSW |
| 10     | Individual 10 | Australian-European | JME    | 918+5G->A             | No           | Female 48y              | 14y                   | Migraine with Aura, PNES, depression | Normal | VPA+LEV | None | Mild ID | GSW |

**Table 1:** Clinical features of individuals with disease-associated variants in KCNQ5.

CAE, childhood absence epilepsy; JME, juvenile myoclonic epilepsy; IGE, idiopathic generalized epilepsy; GTCS, generalized tonic clonic seizure; LOF, loss-of-function; LTG, lamotrigine; ID, intellectual disability; VPA, valproic acid; ETX, ethosuximide; LEV, levetiracetam; TPM, topiramate; ZNS, Zonisamide; PNES, psychogenic non-epileptic seizures; GSW, generalized spike-waves; PPR, photoparoxysmal response; phs, photostimulation; SW, spike-wave; irreg., irregular; GED, generalized epileptiform discharges; PSW, polyspike-and-slow-wave; n/a, not available.
Figure 2. Functional effects of Kv7.5 WT and mutant channels in Chinese hamster ovarian cells. (A) Representative K+ current traces from KCNQ5 WT (black), R359C (green), L692V (orange), Q735R (blue) and untransfected control cells (CTRL, yellow) during voltage steps from -80 mV to +60 mV in 10 mV increments. (B) Peak K+ currents of cells either transfected with WT or one mutated channel subunit were normalized by cell capacitances and plotted versus voltage. All variants result in a significant reduction in current density compared to the WT. WT, n = 11; R359C, n = 10; L692V, n = 10; Q735R, n = 10; CTRL, n = 10. (C) Comparison of maximum peak current density at +60 mV. All variants show a significant reduction compared to the WT. (D) Voltage-dependent activation curves. Lines represent Boltzmann functions fit to the normalized tail current. Currents in the R359C variant were too small to establish such a relationship. (E) Peak K+ currents normalized by cell capacitances and plotted versus voltage of cells either transfected with WT (1 µg) or WT and one mutated channel subunit (1 µg + 1 µg). The significant reduction persisted in all variants compared to the WT indicating a dominant negative effect of the variants on the WT and the significant...
Standard whole-cell patch-clamp recordings were performed from transfected cells which were identified via fluorescent markers localising in the cytosol as they are cleaved off from the subunit by a P2A cleavage site to not affect channel function. Figure 2A displays representative raw current traces recorded from cells expressing either WT or one of the mutant channel subunits. Untransfected cells were used as additional controls. Homomorphic expression of all three mutated channel subunits caused a significant reduction in peak current amplitude and current density (Figure 2B). The R359C variant presented the most severe reduction being almost indistinguishable from untransfected control cells (peaks at 25.2 ± 5.9 pA/pF and 75.5 ± 8.4 pA/pF, respectively; both n = 10; see Table 2). L692V and Q735R reached comparable current densities (112 ± 32 pA/pF and 139 ± 21.4 pA/pF, respectively; both n = 10), of about 20 % of the WT (620 ± 133.3 pA/pF; n = 11; Figure 2C). The voltage dependence of activation, as derived from normalized tail currents, was not changed for L692V and Q735R, and could not be evaluated for R359C, since the currents exhibited by channels carrying this variant were too small to be evaluated (Figure 2D and Table 2).

Homomorphic expression of mutant and WT subunits in a 1:1 ratio did increase the current density for all variants compared to homomorphic expression (Figure 2E). We observed a dominant negative effect for all three variants which was most severe for R359C (peak current density of 49.1 ± 10.7 pA/pF, n = 10) corresponding to 10 % of the WT amplitude (487 ± 54.6 pA/pF; n = 12), while L692V and Q735R reached less than 45 % of the WT (210.7 ± 43.5 pA/pF and 217.5 ± 61.9 pA/pF, respectively; both n = 10) (Figure 2F). The voltage dependence of channel activation was again similar for L692V/WT, Q735R/WT and WT alone, while R359C/WT-associated amplitudes were still too small for data evaluation (Figure 2G).

Since Kv7.5 and Kv7.3 subunits can form heterotetramers,1,2 heteromorphic co-expression of Kv7.5 WT and mutant subunits in a 1:1 ratio (1 μg of each clone) was performed in a CHO cell line stably expressing Kv7.3 WT channels. Antibiotics were removed from the medium 72 hours prior to recordings to ensure comparability with the previous recordings that were conducted in cells cultured in antibiotics-free medium. Current densities were still significantly reduced for all three investigated variants indicating dominant-negative effects also under these conditions. Peaks reached 24 % of the WT (45.1 ± 44.0 pA/pF; n = 13) for R359C (106.8 ± 32.5 pA/pF; n = 10), 38 % for L692V (171.5 ± 31.5 pA/pF; n = 10), and 50 % for Q735R (226.3 ± 66.7 pA/pF; n = 10; Figure 2H and I). Activation curves of all three variants did not significantly differ from the WT (Figure 2).

In a second series of experiments, the effect of the truncating variant A301Gfs*64 (corresponding to c.901dupC) and two additional missense variants (F165I and L926S) were investigated. As these cells were cultured under different conditions using antibiotics in the culture medium (see Materials and Methods), the results of these experiments are shown separately as a second cohort (+ antibiotics), since antibiotics can decrease current density (see Table S1 and Figure S1). Homomorphic expression caused a complete LOF (current density 8.35 ± 1.91 pA/pF; n = 16; Figure 3A to C) compared to the WT (100.0 ± 7.65 pA/pF; n = 69), similar to the effect of the R359C. When co-expressed with WT subunits, neither current density nor activation curves were significantly different in cells expressing the WT subunit alone versus cells expressing WT and variant (see Figure 3D to F and Table 2). Consequently, WT and variant were not co-expressed in the cell line stably expressing Kv7.3, due to the missing effect of the variant on the Kv7.5 WT caused by the absence of the C-terminus due to the variant, and hence, its presumed inability to form heteromers with the WT channels. Both additional missense variants did not show a significant difference in either current density or gating parameters as compared to the WT (see Table 2).

In summary, two missense variants were found to have no functional effect and were thus considered benign, while three other missense variants cause a dominant-negative LOF effect by reducing current density in all three expression conditions, the R359C being the most severe one. The truncated variant (A301Gfs*64) has no effect on the WT subunits, only causing a haploinsufficiency. The voltage dependence of channel activation was not significantly changed for any of the variants (Table 2).

Protein production and membrane expression of KCNQ5 variants in CHO cells

To investigate whether the LOF was caused by a dysfunction in channel opening, or by a trafficking or other defect, the amount of produced protein of the R359C, L692V and Q735R variants compared to the WT was determined in CHO cells via Western blot. The A301Gfs*64 variant had to be excluded from this approach due to the missing C-terminus, which carries the antibody epitope. When whole cell lysates were

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Table 2 provides exact values and statistical analyses.
blotted, no significant difference in protein amount was detected as compared to the WT (Figure 4A and B; n = 3). In addition, expression levels of Kv7.3 and Kv7.5 subunits in the stable Kv7.3 cell line were analysed. We observed a reduction in Kv7.5 subunit expression as compared to Kv7.3 (Figure 4C; n = 4), likely explained by less cells expressing the transiently transfected Kv7.5 than the stably expressed Kv7.3 subunits. To further examine if the mutant subunits were integrated in the cell membrane, a cell surface protein biotinylation assay with subsequent Western blot was performed. Again, no significant differences were observed. WT, n = 40; WT+901dupG, n = 27. (E) Comparison of maximum peak current density at +60 mV. No significant differences were observed. (F) Voltage-dependent activation curves. Lines represent Boltzmann functions fit to the normalized tail current. Shown are means ± SEM. (B, D, F). Scatter-and-whisker plots (C and E) show median (horizontal line) and the interquartile ranges. Dots indicate maximum values of single cells. **** p ≤ 0.0001; ns non-significant; Table 2 provides exact values and statistical analyses.

The R359C variant alters PI(4,5)P2 interaction

R359 is homologous to R360 in Kv7.1, which has been previously described as one of the key PI(4,5)P2 interaction sites.30-31 We performed homology modelling to investigate whether the dominant-negative LOF of the R359C variant in current density could be caused by a change in the interaction of Kv7.5 channels with PI(4,5)P2. Homology models of Kv7.5 and Kv7.5-R359C in absence and presence of PIP3 were generated based on high homology to Kv7.4. Two all-atoms-mobile MD
Table 2: Biophysical properties of $K_{v_{7.5}}$ WT and variant channels.

| Subunit set | Current Density (pA/pF) | Activation kinetics |
|-------------|-------------------------|---------------------|
|             | $V_{1/2}$ (mV) | $k$ |
| WT | 4.29±1.13 | 11 |
| L692V | 3.78±1.66 | 11 |
| L322V | 2.71±1.34 | 13 |
| WT | 4.39±1.30 | 11 |
| R359C | 4.79±1.66 | 13 |

Note: *p* < 0.05 via Kruskal-Wallis with post hoc correction for multiple comparisons with Benjamini, Krugier and Yekutieli's test.

Furthermore, protein-phospholipid overlay assays were performed to investigate whether we could identify altered PIP$_3$ interactions in vitro as well. The assays showed a significant reduction in the binding affinity of R359C channels compared to the WT to phosphatidylinositol 3-phospholipid (PI(3)P; *p* = 0.012), phosphatidylinositol 4-phospholipid (PI(4)P; *p* = 0.002), and most importantly, phosphatidylinositol 3,4-bisphospholipid (PI(3,4)P$_2$; *p* < 0.001), phosphatidylinositol 4,5-bisphospholipid (PI(4,5)P$_2$; *p* = 0.006), and phosphatidylinositol 3,5-bisphospholipid (PI(3,5)P$_2$; *p* = 0.0012; Figure 4F and G; *n* = 3). Our results suggest that R359C plays an important role to confer binding to PI(4,5)P$_2$ that is essential for channel opening, as Thomas et al. (2011) found for the homologous site in KCSQ2.

To elaborate these findings, additional electrophysiological experiments were conducted. To this end, endogenous PIP$_3$ levels were increased by co-expression of a 1y PI(4)P-kinase as reported previously. As a result, WT amplitude and current density significantly increased (to 1182±74±129±13 pA/pF; *n* = 10; see Figure 6A, B, C) as previously reported. Interestingly, the homomeric R359C channels firstly displayed measurable amplitudes under PIP$_3$ co-expression (in 6/10 cells), however, peak current density was still significantly decreased (103±13 ± 98±7 pA/pF) as compared to the WT, only reaching 73±8 pA/pF. Furthermore, the activation curve of R359C channels was significantly shifted towards more depolarized potentials with a $V_{1/2}$ of 7±2±8±2 mV (n = 6; Figure 6D) as compared to WT channels (27±19 ± 5±85 mV; *n* = 10), whereas slopes did not appear to be significantly different (see Table 3). Co-expression of PIP$_3$K did not change activation curve parameters in WT cells as compared to homomeric expression alone. Cells only
expressing PIP5K did not show measurable amplitudes and their peak current density (14.60 ± 3.35 pA/pF; n = 10) did not significantly differ from untransfected CHO cells. These results support the hypothesis that the R359 site might be important for PIP2 interaction and thus channel opening.

To further clarify a potentially decreased PIP2 sensitivity of this variant, additional electrophysiological recordings were conducted by co-expressing a Daino rerio voltage-sensing phosphatase (Dr-VSP) as described by Hossain et al. (2008), which is activated by strong depolarizations (≥ +100 mV) and temporarily reduces PIP2 levels in the cell membrane resulting in a temporary inhibition of Kv7 channels.35 As the R359C variant suppresses currents even under WT co-expression almost completely, the stable Kv7.3-celline was used for these experiments, in which the R359C variant yielded measurable current amplitudes in heterozygous conditions co-expressed with Kv7.5-WT, to be able to see a potential current inhibition by VSP. Activation of VSP for only 0.2 s had an immediate effect on cells transfected with Kv7.5-WT+Kv7.5-R359C+VSP reducing currents by 31 %, whereas cells transfected with Kv7.5-WT+VSP showed an effect only after 0.6 s of VSP activation (n = 5 for all conditions; see Figure 7B) at which the current of 3/5 cells transfected with the variant does not recover from VSP activation anymore. This is further displayed in the current recovery time after VSP switch-
off. While Kv7.5-WT + VSP cells only take 3.2 ± 1.5 s to recover after 0.6 s of VSP activation and 11.8 ± 4.33 s after 1-s (see Figure 7A, C), Kv7.5-WT + Kv7.5-R359C + VSP expressing cells are significantly slowed down (19.0 ± 4.29 s; p < 0.01) and none of the cells recovered to the baseline value, respectively. Co-expression of PIP3K led to a quicker recovery in Kv7.5-WT + VSP cells (1.4 ± 0.75 s and 2.6 ± 1.66 s, respectively) and was able to enhance Kv7.5-WT + Kv7.5-R359C + VSP cells back to normal values (6.4 ± 3.91 s and 13.4 ± 5.33 s, respectively). Consequently, the data strongly suggest that a reduction in PIP3 binding affinity causes the LOF in the R359C variant.

Discussion
Here we analysed clinical and genetic data from multiple cohorts of altogether 1,292 independent families with GGE to identify and functionally characterize five likely disease-related variants suggesting that either haploinsufficiency or dominant-negative effects of KCNQ5 are associated with GGE with or without mild to moderate ID. A hypothesis-driven statistical evaluation in three large GGE cohorts and matched controls indicated that rare, functionally relevant variants in KCNQ5 might be more frequent in GGE than expected by chance. Since KCNQ5 was not among the top-ranking genes in the exome-wide primary analyses of these studies, it appears that pathogenic variants in KCNQ5 are rare.

The phenotypes ranged from mild CAE to pharmaco-resistant early-onset absence epilepsy or JME with moderate ID. Most of the individuals (8/10) presented with absence seizures. In four individuals, developmental delay or ID was present prior to epilepsy. The phenotype comprising absence seizures and ID is in
accordance with a previous report of an individual with mild ID and absence seizures in adolescence, carrying an intragenic duplication in KCNQ5 leading to haplo-insufficiency.19 Lehman and colleagues reported four individuals suffering from ID and/or epilepsy caused by KCNQ5 variants, which they described as LOF and one GOF (P369R) carried by the most severely affected individual. However, a recent study showed that these variants along with four additional missense variants found in children with mild to severe ID and epilepsy in fact caused a GOF in channel activation and deactivation. Two additional truncating variants including a LOF were found in individuals with a milder phenotype. Moreover, this study shows that the more severe GOF variants were found in more severely affected individuals, while milder GOF variants and the truncating LOF variants belonged to individuals with a milder phenotype.17 This is in line with a second recent publication that found two pore variants to cause a GOF in individuals with DEE.18 The individuals presented here show a milder phenotype as compared to the three studies mentioned and two carriers were even asymptomatic (one with R359C and one with Q735R). While global developmental delay of varying degree is a common feature in all described individuals in the above mentioned studies, three of our variants were not associated with ID (L692V, Q735R and A301Gfs*64) and the family carrying the R359C variant shows a spectrum with one individual being unaffected to individuals with mild to moderate ID. Interestingly, all four individuals from Lehman et al. suffered from ataxia with a varying degree of severity. In contrast, none of the individuals described by Wei et al. and Nappi et al. had ataxia and only one individual of our cohort, the most severely affected one, had mild ataxia on neurological examination, the others did not have neurological abnormalities on examination. Moreover, all of the individuals presented here presented with generalized seizures, yet not all of the individuals described in the other studies suffer from seizures. Five of ten of the individuals presented here presented with generalized seizures, yet not all of the individuals described in the other studies suffer from seizures. Five of ten of the individuals presented here presented with generalized seizures, yet not all of the individuals described in the other studies suffer from seizures. Five of ten of the individuals presented here presented with generalized seizures, yet not all of the individuals described in the other studies suffer from seizures. 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(B) Peak K+ currents of cells either transfected with WT alone (grey dots), WT + PIP5K (black), R359C (grey triangles) or R359C + PIP5K (green triangle) channel subunit were normalized by cell capacitances and plotted versus voltage. The currents generated by KcNQ5 WT + PIP5K remain largely reduced as compared to KcNQ5 WT + PIP5K. KcNQ5 WT, n = 12; KcNQ5 WT + PIP5K, n = 10; KcNQ5-R359C, n = 10; KcNQ5-R359C + PIP5K, n = 10; PIP5K, n = 10; CTRL n = 10. (C) Comparison of maximum peak current density at +60 mV. PIP5K co-expression significantly increased the KcNQ5-R359C peak current density, yet it is still significantly reduced as compared to KcNQ5 WT + PIP5K. (D) Voltage-dependent activation curves. Lines represent Boltzmann functions fit to the normalized tail current. The activation curve for KcNQ5-R359C + PIP5K is significantly shifted towards more positive voltages. Shown are means ± SEM (B, D). Scatter-and-whisker plots (C) show median (horizontal line) and the interquartile ranges. Dots indicate maximum values of single cells. **** p ≤ 0.001; **p ≤ 0.01; * p ≤ 0.05; ns non-significant; Table 3 provides exact values and statistical analyses.
The results of our functional analysis show a LOF in current density for homomeric channels for all investigated variants, and three of the investigated variants have a dominant-negative effect on current density in heteromeric expression experiments with a near complete loss-of-function for the R359C variant. The frameshift variant (A301Gfs*64) results in a premature stop codon in the pore region deleting the entire C-terminus. As the C-terminus comprises the interaction sites for the subunits to form a channel, its absence causes this variant to be unable to form channels under homomeric expression and abolishes interaction with the WT subunits under heteromeric conditions causing a haploinsufficiency, but not a dominant-negative effect on the WT subunits. According to our electrophysiological studies, members of family 1 carried the most severe variant (R359C), leading to a severe LOF with a dominant-negative effect. We could not find clear genotype-phenotype correlations within our study, since (i) family 1 carrying the most severe variant displays a large phenotypic heterogeneity ranging from mildly to severely affected, pharmaco-resistant individuals, (ii) individuals in family 3 exhibited a similarly severe epileptic phenotype albeit the Q735R variant showed a less prominent electrophysiological dysfunction, and (iii) also the phenotype of previously reported individuals carrying variants which only caused haploinsufficiency were reported with similar or more severe phenotypes.16,17,18,19 This is in contrast to other Kv7 channels, in which functionally more severe variants with dominant-negative effects cause more severe epileptic phenotypes.35 Looking at our results in the context of the newly published variants, it seems that individuals carrying LOF variants display milder phenotypes, while GOF variants cause more severe phenotypes, such as DEE. Additionally, the functional severity of the GOF seems to correlate with the severity in the phenotype of the patient.17 Similar patterns of GOF variants inducing much more severe phenotypes such as DEE and LOF variants inducing milder phenotypes such as GGE have been described for other genes such as SCN8A,16 SCN2A17 and KCNA2.38 For KCNQ4, it seems that any significant LOF can contribute to an epileptic phenotype or ID of varying severity, which might be influenced by other individually differing factors, such as compensatory effects, the genetic background or environmental determinants. Larger cohorts are needed to further investigate this issue.

Remarkably, all missense variants showed stable total and membrane-expressed protein levels in CHO cells as compared to the WT. The LOF in current density is thus not caused by defects prior to membrane insertion of the channel, such as an abolished tetramersization or trafficking defect, as have been described for variants in KCNQ2 and KCNQ3.15,19 Rather, the three investigated missense variants have functional effects on channel gating, and thus, might mark important sites involved in channel opening. Kv7 channels form
the molecular basis of M-currents and are classically negatively regulated by muscarinic acetylcholine receptors via a PI(4,5)P2-dependent mechanism. PI(4,5)P2 is required for the stabilization of the open state relative to the closed state and PI(4,5)P2-depletion upon activation of muscarinic acetylcholine receptors leads to channel closure. Here, we provide modelling, electrophysiological and biochemical evidence that the R359C variant causes altered PI(4,5)P2 binding, which possibly explains the complete, dominant-negative LOF caused by this variant.

Both other variants (L692V and Q735R) that we investigated are located in an evolutionary highly conserved region of the Kv7.5 C-terminus, which does not show any variants in unaffected individuals in multiple databases and is absent or not conserved in other Kv7 channels, underlining the importance of this region for proper channel function in Kv7.5. As the functional aspects of this region on channel behaviour have not been described previously, these two variants might be able to elucidate binding partners and disclose the function of the distal C-terminus in channel opening. Investigating these molecular mechanisms may open doors for new treatment options in the future, especially for pharmaco-resistant patients.

In summary, we have identified rare loss-of-function variants in KCNQ5 in five independent families, which are likely contributing to the pathophysiology of GGE. Two variants cause haploinsufficiency, three showed a dominant-negative effect on WT KV7.5 and KV7.3 channels. We were also able to identify the importance of R359 as crucial for PI(4,5)P2 interaction and channel opening. Consequently, the M-current in these individuals is likely reduced causing a decrease in action potential threshold and increased excitability of neurons expressing Kv7.5 channels, thus leading to an elevated seizure susceptibility. The types of neurons and networks that are involved need to be determined in further
studies. Furthermore, identifying these LOF variants in these patients opens doors to targeted treatment using Kv7 channel openers such as retigabine, and further studies should be conducted to investigate their effect on the variants.

Contributors
JKr, JS, PM, GL, SM and HL designed the study and experiments. JKr, JS, AL, JJH, MM, GS, PY, MK, MSH, PM performed experiments and analyzed data. JKr, JS, AL, JJH, MM, GS, PY, MK, SP, RK, PM, GL, SM, and HL, interpreted data. JKe, KA-K, HC, BJS, YGW, PK-K, SFB, GL, and HL recruited and phenotyped patients. JKr, JKe, GS, MK, PM, GL, SM, and HL wrote the manuscript. All authors read, revised and approved the manuscript.

Data sharing statement
The exome sequencing data/analyses presented here are based on the use of study data from the Epi25 Collaborative (http://epi-25.org/), available with controlled access through dbGaP (https://ncbi.nlm.nih.gov/gap/), the EuroEPINOMICS-CoGIE project, and the Epi4K project.

Declaration of interests
J. Krüger was financed by a grant from the Deutsche Forschungsgemeinschaft (DFG), during the conduct of the study; Dr. Schubert has nothing to disclose; Dr. Kegele has nothing to disclose; A. Labalme has nothing to disclose; Dr. Mao has nothing to disclose; J. Heighway has nothing to disclose; Dr. Seebohm has nothing to disclose; Dr. Yan has nothing to disclose; M. Koko reports grants from DAAD, outside the submitted work; Dr. Aslan has noth-thing to disclose; Dr. Caglayan has nothing to disclose; Dr. Steinhoff has nothing to disclose; Dr. Weber has nothing to disclose; Dr. Keo Kosal has nothing to disclose; Dr. Berkovic reports grants from NHMRC, during the conduct of the study; grants from UCB Pharma, grants from Eisai, grants from SciGen, personal fees from Bionomics, personal fees from Athena Diagnostics, outside the submitted work; In addition, Dr. Berkovic has a patent Methods of treatment, and diagnosis of epilepsy by detecting mutations in the SCN1A gene with royalties paid to Patent held by Bionomics Inc. Licensed to Athena Diagnostics; Genetics Technologies Ltd, a patent Diagnostic and Therapeutic Methods for EFMR (Epilepsy and Mental Retardation Limited to Females) with royalties paid to Licensed to Athena Diagnostics, and a patent A gene and mutations thereof associated with seizure and movement disorders (PRRT2) with royalties paid to Licensed to Athena Diagnostics; Dr. Hildebrand has nothing to disclose; Dr. Petrou reports personal fees and other from Praxis Precision Medicines, outside the submitted work; and Dr. Petrou works for a company, Praxis Precision Medicines that develop therapies for neurogenetic disorders such as KCNQ5 (but this is not currently under any consideration); Drs. Krause and May have report grants from the Fondation de la Recherche en Luxembour; Dr. Lesca has nothing to disclose; Dr. Maljevic has nothing to disclose; Dr. Lerche reports grants from the German Research Foundation (DFG), from the Federal Ministry for Education and Research (BMBF), grants from Foundation no epilep, during the conduct of the study; outside the submitted work, Dr. Lerche reports a grant from the Else-Kröner Fresenius Foundation (EKFS), a grant and personal fees from Bial, a grant from Boehringer Ingelheim, personal fees from Eisai, personal fees from UCB/Zogenix, personal fees from Arvelle/Angelini Pharma, personal fees from Desitin, and personal fees from IntraBio.

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Supplementary materials
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