The study of sodium and potassium channel gene single-nucleotide variation significance in non-mechanical forms of epilepsy

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Abstract

Background: Epilepsy is one of the most common and heterogeneous neurological diseases. The main clinical signs of the disease are repeated symptomatic or idiopathic epileptic seizures of both convulsive and non-convulsive nature that develop against a background of lost or preserved consciousness. The genetic component plays a large role in the etiology of idiopathic forms of epilepsy. The study of the molecular genetic basis of neurological disorders has led to a rapidly growing number of gene mutations known to be involved in hereditary ion channel dysfunction. The aim of this research was to evaluate the involvement of single-nucleotide variants that modify the function of genes (SCN1A, KCNT1, KCNT1, and KCNQ2) encoding sodium and potassium ion channel polypeptides in the development of epilepsy.

Results: De novo mutations in the sodium channel gene SCN1A c.5347G>A (p. Ala1783Thr) were detected in two patients with Dravet syndrome, with a deletion in exon 26 found in one. Three de novo mutations in the potassium channel gene KCNT1 c.2800G>A (p. Ala934Thr), were observed in two patients with temporal lobe epilepsy (TLE) and one patient with residual encephalopathy. Moreover, a control cohort matched to the case cohort did not reveal any SNVs among conditionally healthy individuals, supporting the pathogenic significance of the studied SNVs.

Conclusion: Our results are supported by literature data showing that the sodium ion channel gene SCN1A c.5347G>A mutation may be involved in the pathogenesis of Dravet syndrome. We also note that the c.2800G>A mutation in the potassium channel gene KCNT1 can cause not only autosomal dominant nocturnal frontal lobe epilepsy (ADNFLE) but also other forms of epilepsy. To treat pathogenetic mutations that accelerate the function of sodium and potassium ion channels, we recommend ion channel blockade drug therapy.

Keywords: Epilepsy, Mutation, De novo, Ion channel gene

Background

Epilepsy is one of the most common serious neurological disorders, affecting 4–10 per 1000 people and approximately 50 million people worldwide [1]. Epilepsy comprises a group of chronic brain diseases characterized by chronic recurrent unprovoked seizures caused by abnormal excessive electrical discharges of brain neurons [2]. Epilepsy can be induced by both mechanical and non-mechanical factors. Approximately 20–30% of epilepsy cases are caused by acquired conditions such as stroke, swelling, or head trauma. Non-mechanical epilepsy forms are associated with hereditary predisposition. Recent data indicate a genetic background in 70–80% of epilepsy cases [3, 4]. Moreover, some hereditary syndromes (Rett, Dravet, Angelman, West, Prader-Willi, and others) are accompanied by epileptic seizures.

Molecular genetic studies of neurological disorders have led to a rapidly growing number of gene mutations known to be involved in hereditary ion channel dysfunction. The study of the molecular genetic basis of neurological disorders has led to a rapidly growing number of gene mutations known to be involved in hereditary ion channel dysfunction. The aim of this research was to evaluate the involvement of single-nucleotide variants that modify the function of genes (SCN1A, KCNT1, KCNT1, and KCNQ2) encoding sodium and potassium ion channel polypeptides in the development of epilepsy.
dysfunction (genetic channelopathies). The normal functioning of ion channels is especially important in the nervous system for the generation, repression and distribution of action potentials [5, 6]. Taking into account their importance in neuronal excitability and synaptic transmission through the central and peripheral nervous systems, it is not surprising that mutation of the genes responsible for ion channel functioning can cause epilepsy. Most hereditary epilepsy forms with detected gene mutations are caused by changes to ion channels that ensure neuronal membrane polarization. They include genes encoding sodium, potassium, calcium, and chloride channels (SCN1A, SCN2A, CACNA1A, KCN10, KCNT1, KCN1C, KCNQ2, CLCN1) [3, 4, 7–10].

Nonetheless, the list of candidate ion channel genes involved in epilepsy is not restricted by variants in the above genes. Due to large-scale genome-wide studies (GWASs) of epilepsy patients, the spectrum and number of gene mutations possibly involved in epilepsy pathogenesis are increasing every year [11]. However, the low frequency of mutation requires more detailed screening of certain single-nucleotide variants (SNVs) in healthy individuals and in those with different forms of epilepsy.

The aim of this research was to examine involvement in the development of epilepsy single-nucleotide variants that modify the function of genes encoding the polypeptides of sodium and potassium ion channels, as follows: SCN1A (rs121918748, c.5459T>C-p.Phe1820Ser [12], rs571447839, c.5347G>A-p.Ala1783Thr [13], rs121918792, c.5020G>C-p.Gly1674Arg [14], rs121917948, c.4969C>T-p.Pro1657Ala [15], KCNT1 (rs397515405, c.2782C>T-p.Arg928Cys [16] and rs397515403, c.2800G>A-p.Ala934Thr [17]); KCNQ1 (rs727502818, c.959G>A-p.Arg320His [12]); KCNQ2 (rs587777219, c.794C>T-p.Ala265Val [18] and rs28939683, c.851A>G-p.Tyr284Cys [19]).

Methods

Sampling

This “case-control” study was approved by the Local Ethics Committee of Kazakh-Russian Medical University (protocol N.51, 05.09.2017). The studied cohorts included 120 patients with different forms of epilepsy of non-mechanical origin and 120 conditionally healthy individuals. In some cases, the patient’s first-degree relatives were included (17 persons). Peripheral blood samples were collected on the basis of the SVS clinic by V.M. Savinov (Almaty, Kazakhstan) and Center of Neurology, Epileptology, and EEG, “Arnur” (Almaty, Kazakhstan). Detailed questionnaires and informed consent were obtained when samples were collected. Epilepsy diagnosis was based on clinical symptoms, electroencephalograms (EEG), and magnetic resonance imaging (MRI) data.

Genotyping of SNVs

DNA samples were isolated using Genefet Genomic DNA Purification Kit (Thermo Fisher Scientific, USA) in accordance with the protocol recommended by the manufacturer.

Site-specific PCR amplification with restriction of amplified fragments (PCR-RFLP) was used for genotyping ion channel gene SNVs (rs121918748, rs571447839, rs121918792, rs121917948, rs397515405, rs397515403, rs727502818, rs137989254, rs587777219, rs28939683). Primers were designed by using the PrimerQuest Tool online program. The PCR mixture contained 20–50 ng genomic DNA, 10 pM each specific primer, and 10 μL Master mix (2x, Thermo Fisher Scientific, USA). Temperature and time conditions were selected for each SNV by taking into account the size of the primers used (Table 1). Genotyping of SNVs was performed by PCR with subsequent restriction fragment length polymorphism (RFLP) analysis in accordance with the protocol recommended by the manufacturer (Thermo Fisher Scientific, USA).

Restriction endonucleases were selected with the help of WatCut for SNP-RFLP online analysis. The RFLP details are represented in Table 2.

Statistical analysis

The standard chi² test was used to assess the significance (P values) of the observed differences between the case and control groups. An alpha error (P) of less than 0.05 was used as the criterion for significance.

Allele frequencies were calculated in accordance with standard Hardy-Weinberg equilibrium.

Results

Characteristics of the studied cohorts

Clinical examination of epilepsy patients confirmed the non-mechanical nature of the different epilepsy forms in the individuals in the study cohort. In all patients, neurological status did not show meningeal signs or cerebral symptoms. A memory/attention decrease and emotional liability in the psycho-emotional sphere were detected in one patient. In total, 71 patients had a pathological EEG; 10 individuals had a normal EEG. MRI detected pathologies for 49 individuals, whereas 12 did not have pathological signs. Seizure manifestations allowed us to diagnose the following epilepsy types: temporal epilepsy—10 cases; symptomatic epilepsy—20 cases; residual encephalopathy—14 cases; frontal epilepsy—4 cases; idiopathic epilepsy—18 cases; juvenile myoclonic epilepsy—6 cases; absence epilepsy—3 cases; myoclonic epilepsy—case; West syndrome—1 case; Angelman syndrome—2 cases; Dravet syndrome—3 cases; Rett syndrome—1 case; Lennox Gastaut syndrome—1 case. For
other patients, the exact type of epilepsy was not determined.

The control cohort representing people without any noticeable neurological pathologies was matched to the case cohort by age, sex, and ethnicity (Table 3).

**SNV analysis in the control and case cohorts**

The selected ion channel gene SNVs (*SCN1A* 26 exon c.5459T>C, c.5347G>A, c.5020G>C, and c.4969C>T; *KCNT1* 24 exon c.2782C>T, and c.2800G>A; *KCNQ2* 5 and 6 exon c.794 C>T; *KCNQ1* 2 exon c.959 G>A; *KCNJ1* 5 and 6 exon c.794 C>T, and c.851 A>G) were genotyped in the control and case cohorts.

The studied SNVs were not detected in the control cohort of conditionally healthy persons. Two patients with Dravet syndrome carried mutations in the sodium channel gene *SCN1A*. Figure 1 demonstrates the PCR-RFLP results for *SCN1A* c.5347 G>A (p.Ala1783Thr) mutation screening.

The *SCN1A* c.5347 G>A (p.Ala1783Thr) mutation was detected in a heterozygous state in an 8-year-old patient. Convulsions first appeared at the age of 3 months and were repeated 2 times a month with different semiotics. Febrile convulsions were not registered. Valproate treatment led only to a slight improvement; treatment was replaced by topiramate, which resulted in a decrease in seizure frequency; worsening of the patient’s condition was observed after oxcarbazepine treatment. Based on this, treatment with topiramate in combination with valproate and dexamethasone was recommended. However,

| Gene, location | SNVs | Primers, 5′→3′ |
|---------------|------|----------------|
| SCN1A, 26 exon | c.5499T>C (p.Phe1820Ser) | f-CCCGACTGACACCTGTTTTTTAAATAGAG GTCCTGTTTTTTAAATAGAG |
| | c.5347G>A (p.Ala1783Thr) | r-GTAGGTGTTTTTTAAATAGAG GTCCCTTCTGTTTTTTAAATAGAG |
| | c.5020G>C (p.Gly1674Arg) | r-CTGACCAAACCTTCTCCTTCTT |
| | c.4969C>T (p.Pro1657His) | r-GTCCCTGACACCTTCTCCTTCTT |
| KCNT1, 24 exon | c.2782C>T (p.Arg928Cys) | r-GTAGGTGTTTTTTAAATAGAG GTCCCTTCTGTTTTTTAAATAGAG |
| | c.2800G>A (p.Ala934Thr) | r-CTGACCAAACCTTCTCCTTCTT |
| KCNQ2, 5 and 6 exon | c.794 C>T (p.Ala265Val) | r-GTAGGTGTTTTTTAAATAGAG GTCCCTTCTGTTTTTTAAATAGAG |
| | c.851 A>G (p.Tyr284Cys) | r-GTAGGTGTTTTTTAAATAGAG GTCCCTTCTGTTTTTTAAATAGAG |
| KCNC1, 2 exon | c.959 G>A (p.Arg320His) | r-GTAGGTGTTTTTTAAATAGAG GTCCCTTCTGTTTTTTAAATAGAG |

Table 1: The site-specific PCR amplification protocols

| Gene, location | Mutation/polymorphism/primers (3′→5′) | Restriction endonuclease | DNA fragments length and corresponding genotype |
|---------------|----------------------------------------|--------------------------|-----------------------------------------------|
| SCN1A, 26 exon | c.5459T>C (p.Phe1820Ser) | PstI | TT-321 bp; CC-282 and 39 bp; TC-282, 39 and 282 bp |
| | c.5347G>A (p.Ala1783Thr) | Acc II | GG-188 and 133 bp; AA-321 bp; GA-321, 188 and 133 bp |
| | c.5020G>C (p.Gly1674Arg) | HaeIII | GG-90, 83 and 75 bp; CC-173 and 75 bp; GC-173, 90, 83 and 75 bp |
| | c.4969C>T (p.Pro1657His) | BanHI | CC-140 and 108 bp; TT-248 bp; CT-248, 140 and 108 bp |
| KCNT1, 24 exon | c.2782C>T (p.Arg928Cys) | HpyF10VI | CC-116 and 117 bp; TT-233 bp; CT-233, 117 and 116 bp |
| | c.2800G>A (p.Ala934Thr) | Acc II | GG-128 and 105 bp; AA-233 bp; GA-233, 128 and 105 bp |
| KCNQ2 5 and 6 exons | c.794 C>T (p.Ala265Val) | RsaI | CC-175 and 35 bp; TT-210 bp; CT-210, 175 and 35 bp |
| | c.851 A>G (p.Tyr284Cys) | AccII | AA-156 and 65 bp; GG-221 bp; AG-221, 156 and 65 bp |
| KCNC1 2 exon | c.959 G>A (p.Arg320His) | HpyCH4V | GG-519 bp, AA-367 and 152 bp; GA-519, 367 and 152 bp |

Table 2: Restriction endonucleases and SNV specific RFLP details
despite this, convulsions occurred daily with myoclonia of the eyes and shoulders.

The deletion (33 bp) of exon 26 of the *SCN1A* gene was detected in another patient, a 4-year-old child (Fig. 2). The first seizures of this patient were marked by twitching of the right hand with gradual involvement of the leg and secondary generalization; at present, attacks begin with tonic tension with subsequent clonic twitches. The EEG for sleeping time demonstrated sharp evoked potentials in the adductions of the left posterior temporal domain and series of bitemporal asynchronous theta waves. As the *SCN1A* gene deletion was in a heterozygous state, the presence of a second functional copy provides partial *SCN1A* gene function preservation. Despite the identified mutation, the patient's response to treatment was adequate.

The c.2800G>A (p.Ala934Thr) mutation of the potassium ion channel gene *KCNT1* was detected in 3 patients (Fig. 3); two of these patients had temporal lobe epilepsy, and one patient had residual encephalopathy.

Both patients with temporal lobe epilepsy (years of birth 1972 and 1988) had psychomotor automatism attacks with partial seizures. MRI of the patient born in 1972 detected residual subatrophic changes in the brain, and EEG showed a pathological variant that was more indicative of the temporal lobe. The second patient (born in 1988) first experienced attacks at the age of 20. A computer tomography scan of the brain showed that the patient had mild ventriculomegaly of the lateral ventricles, and EEG revealed polymorphic dysrhythmia slightly increased in the temporal-parietal ventricles. The patient was treated with carbamazepine 600 mg/day, but despite this, attacks were occurred every month.

The third carrier of the *KCNT1* codon 934 mutation was a patient born in 1987 who had primary generalized tonic-clonic seizures that developed suddenly. MRI revealed congenital peculiarity of mid-brain structure development in the background of non-rough changes in residual genesis.

We performed molecular genetic examinations of the first-degree relatives of patients with detected SNVs to determine the hereditary burden. We did not find *SCN1A* exon 26 or *KCNT1* exon 24 mutations among the close relatives of the indicated patients, supporting the de novo origin of the *SCN1A* c.5347G>A and *KCNT1* c.2800G>A mutations detected in the children of healthy parents.

**Discussion**

In Kazakhstan, more than 70,000 people have epilepsy, of which 28,000 are children, adolescents, and young people; 38% of patients become disabled, and their

| Table 3 The main characteristics of studied cohorts |
|--------------------------------------------------|
| **Cohort** (sample volume) | **Years of birth (average age)** | **Sex (%)** | **Ethnicity, persons (%)** |
| Case (120) | 1960–2017 (21.00 ± 13.00) | Men | Women | Kazakhs | Russian | Other |
| 74 (61.67) | 46 (38.33) | 84 (70.00) | 23 (19.17) | 13 (10.83) |
| Control (120) | 1966–2018 (20.72 ± 11.58) | 68 (56.67) | 52 (43.33) | 88 (73.33) | 23 (19.17) | 9 (7.50) |
| t = 0.01608 | 0.50219 | 0.60277 | 0.30476 | 0 | 0.81371 |
| P = 0.98976 | 0.70371 | 0.65466 | 0.81169 | 1 | 0.56516 |

**Fig. 1** PCR-RFLP detection of *SCN1A* gene mutation p.Ala1783Thr (c.5347G>A). M–DNA Ladder GeneRuler 100 bp (Thermo Fisher Scientific, USA); 1,2,5,6,11–homozygotes by normal allele c.5347G>A (188 and 133 bp DNA fragments length); 3 and 7–c.5347G>A heterozygote genotype (321, 188, and 133 bp DNA fragments length)
quality of life decreases by an average of 85%. All patients in this study had generalized idiopathic (27%), focal (47%), and epileptic encephalopathies (26%). The focal seizure type includes symptomatic (76.8%) and idiopathic forms (23.2%) [20].

This study regarding the molecular genetic spectrum of epilepsy-associated causative mutations is the first conducted in Kazakhstan. In general, statistical data about epilepsy hereditary form frequency are unavailable, which was the main reason why we chose non-mechanical epilepsy forms, as they can be associated with genetic disorders. The matched control cohort was selected from conditionally healthy individuals after sampling epilepsy cases, taking into account sex, ethnicity, and age.

Worldwide data on the genetics of epilepsy and hereditary syndromes characterized by epileptic seizures are very limited. Basically, data on complete screening of the genomes of patients with various forms of epilepsy are available [21]. As a rule, all identified genetic changes in ethnically heterogeneous populations are single cases or are characterized by a low frequency (less than 1% of the general population frequency) [22].

Analyzing the available literature sources on the spectrum of genes associated with epilepsy, it is clear that mutations of ion channel genes that alter their function are most often recorded [23–25]. Only a few cases of heterozygous mutations were found in 120 patients with different forms of epilepsy.
Moreover, none of the studied SNVs was detected in the control cohort of conditionally healthy persons, though this situation did not allow us to apply the statistical treatment-odds ratio method (OR). Based on the available comparability of case and control cohorts by age, sex, ethnic background, and cohort volume, we consider that the mutations detected (SCN1A c.5347 G>A (p.Ala1783Thr) and KCNT1 c.2800G>A (p.Ala934Thr) have pathogenic significance. In both cases, replacement of a non-polar aliphatic amino acid alanine with a polar oxyaminocarboxylic threonine occurs, which results in a small physicochemical difference of the corresponding polypeptides of sodium and potassium ion channels. Experimental studies have shown that this missense change renders the KCNT1 ion channel constitutively active when assayed in cell culture [17, 26]. The SCN1A p.Ala1783Thr missense variant was first described in 2007 [27] as being associated with Dravet syndrome, and it is reported as pathogenic in the following databases: (i) HGMD, (ii) Ensembl, and (iii) ClinVar. According to the ACMG criteria, the variant is also classified as pathogenic (PS2, PS3, PM1, PM2, PP2, PP3, PP4, PP5) [28]. This variant is likely to be disruptive, but these predictions have not been confirmed by published functional studies, and their clinical significance is still uncertain. These mutations are proposed to accelerate the functioning of sodium and potassium ion channels [23], causing epileptic seizures.

The frequency of detected SNVs in exon 26 of the sodium channel gene SCN1A (c.5347G>A (p.Ala1783Thr), rs571447839) was 0.01667, and it was 0.00833 for the 33 kb deletion. The frequency of SNVs in exon 24 of the potassium ion channel gene KCNT1 (c.2800G>A (p.Ala934Thr), rs397515403) was recorded as 0.025.

Our results show a more notable frequency for the SNVs rs571447839 and rs39751540 compared to the frequency in global databases (ClinVar, 1000G, Esp6500, ExAC), which can be explained by geographical location or ethnic peculiarities of the studied cohort of epilepsy patients.

Mutations in the SCN1A gene are mostly described for patients with Dravet syndrome, as well as for other epileptic forms [29–32]. Mutations in the SCN1A gene are inherited autosomal dominantly and lead either to a loss or a change in function. For instance, mutations related to loss of function are more likely associated with Dravet syndrome [33]. However, SCN1A mutations altering sodium ion channel function are mostly recorded for generalized epilepsy with febrile seizures [30]; mutations of this gene are de novo in 95% of cases [27]. Two studies [13] recorded single de novo cases of SCN1A c.5347G>A mutation in patients with Dravet syndrome. Our results are in accordance with these data. Based on a study of first-degree relatives of patients carrying the SCN1A c.5347G>A mutation and 33 kb deletion, we concluded the de novo occurrence of these mutations in the studied families.

Autosomal dominant mutations in the potassium channel gene KCNT1 are associated with the development of epileptic syndromes such as MMPSI [16, 17, 34, 35], ADNFLE [16], early infantile epileptic encephalopathy (EIEE), and Ohtahara syndrome (OS) [34, 36]. Heterozygous mutations of the KCNT1 gene have been described for patients with inatlante epilepsy with migratory focal seizures [17], one patient with leukoencephalopathy, and one patient with severe epilepsy [36].

Ohba C et al. detected 9 heterozygous mutations in the KCNT1 gene in 11 patients, 10 of which were described as de novo mutations; 1 patient inherited the KCNT1 mutation from a mother with mosaicism [37]. Interestingly, some mutations of the KCNT1 gene (including the KCNT1 c.2800G>A mutation, which was found in our case cohort) are associated with one of two different phenotypes, ADNFLE or MMFSI, even within the same family [38]. This indicates that the relationships of genotype and phenotype for KCNT1 c.2800G>A mutation can occur in patients with temporal lobe epilepsy (TLE) and residual encephalopathy, which indicates the clinical heterogeneity of genetic disorders associated with the KCNT1 gene.

Conclusion
Our results indicate the possibility of detecting pathological mutations in the SCN1A and KCNT1 genes, even in a small cohort of patients with non-mechanical epilepsy forms, without expensive genome sequencing. Because of high frequency of KCNT1 c.2800G>A (p. Ala934Thr) mutation (0.025) in epilepsy patients from Kazakhstan, we recommend screening for this mutation in patients from the Central Eurasian region with different epilepsy forms, including MMPSI, ADNFLE, EIEE, TLE, and residual encephalopathy. In the future, we plan to use next-generation sequencing (386 genes in the epilepsy panel) to expand the search for new genes and mutations. We hope that this strategy, accompanied primarily by the screening of mutations in other ion channel genes, will help to develop effective therapy protocols for epilepsy patients with non-mechanical forms. We suppose that, in the case of detection of ion channel gene mutations that lead to Na, K channelopathies, treatment by agents limiting the distribution of electric potential will be effective. In the case of detecting pathogenetic mutations that accelerate the function of sodium and potassium ion channels, we recommend using therapy based on the effect of ion channel blockade. For example, valproic acid or sodium valproate can block...
sodium ion channels; phenytoin or carbamazepine block sodium and potassium ion channels.

Abbreviations
SME: Severe myoclonic epilepsy of infancy; ADNFLE: Autosomal dominant night frontal lobe epilepsy; MMP3: Malignant migrating partial seizures of infancy; GS: Ohtahara syndrome; EIEE: Early infantile epileptic encephalopathy; TLE: Temporal lobe epilepsy; GWASSs: Genome-wide studies; SNV: Single-nucleotide variants; EEG: Electroencephalograms; MRI: Magnetic resonance imaging; PCR: Polymerase chain reaction; RFLP: Restriction fragment length polymorphism

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Authors' contributions
DL and OK conceived and designed the study, collected, assembled and interpreted the data, and wrote the manuscript. YE and OA clinically evaluated the patient’s interpreted data and wrote the manuscript. ZT, SM, and GB performed the experiments and helped with writing of the manuscript. AP and NK helped critically in revising the manuscript. All authors read and approved the final manuscript.

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Availability of data and materials
The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Ethics approval and consent to participate
The authors declare that they have no competing interests.

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References
1. World Health Organization. Epilepsy. key facts. https://www.who.int/news-room/fact-sheets/detail/epilepsy. Accessed 20 Aug 2020.
2. Chang BS, Lowenstein DH (2003) Epilepsy. N Engl J Med 349:1257–1266. https://www.nejm.org/doi/full/10.1056/NEJMoa023208
3. Hilbrandt MS, Dahl HH, Damiano JA, Smith RJ, Scheffer IE, Berkovic SF (2013) Recent advances in the molecular genetics of epilepsy. J Med Genet 50:271–299. https://doi.org/10.1136/jmedgenet-2012-101448
4. Wang J, Lin ZJ, Liu L, Xu HQ, Shi YW, Yi YH, He N, Liao WP (2017) De novo SCN1A mutations in patients with SCN1A mutations. Epilepsia 47(10):1629–1635. https://doi.org/10.1111/epi.14163
5. Heron SE, Scheffer IE, Iona X, Zuberi SM, Birch R, McMahon JM, Bruce CM, Berkovic SF, Muley JC (2010) De novo SCN1A mutations in Dravet syndrome and related epileptic encephalopathies are largely of paternal origin. J Med Genet 47:137–141. https://doi.org/10.1136/jmg.2008.065912
6. Shi X, Yasumoto S, Nakagawa E, Fukasawa T, Uchiya S, Hirose S (2009) Missense mutation of the sodium channel gene SCN2A causes Dravet syndrome. Brain Dev 31:758–762. https://doi.org/10.1016/j.braindev.2009.08.009
7. Myers TC, Mefford HC (2015) Advancing epilepsy genetics in the genomic era. Genome Med 7(1):61. https://doi.org/10.1186/1755-8166-7-61
8. Wang J, Lin ZJ, Liu L, Xu HQ, Shi YW, Yi YH, He N, Liao WP (2017) De novo SCN1A mutations in Dravet syndrome and related epileptic encephalopathies are largely of paternal origin. J Med Genet 47:137–141. https://doi.org/10.1136/jmg.2008.065912
9. Poryo M, Clasen O, Oehl-Jaschkowitz B, Christmann A, Gortner L, Meyer S, Younkin PD, Dlugos JD, Petrovski S, Goldstein BD, Dibbens ML, Scheffer EI, Dibbens LM (2012) Missense mutations in the sodium-gated potassium channel gene KCN7 cause severe autosomal dominant nocturnal frontal lobe epilepsy. Nat Genet 44(11):1188–1190. https://doi.org/10.1038/ng.2440
10. Mill M, Lacoste C, Cacciaviglio P et al (2015) Variable clinical expression in patients with mosaicism for KCNQ2 mutations. Am J Med Genet A 167A(10):2314–2318. https://doi.org/10.1002/ajmg.a.37152
11. Vezzani A, Aronica E, Di Russo E (2013) Early-onset epileptic encephalopathy: from genes and mechanisms to disease-targeted therapies. Pharmacol Rev 70(1):142–173. https://doi.org/10.1124/pr.111.110613
12. D’Adamo MC, Liantonio A, Conte E et al (2020) Ion channels involved in neurodevelopmental disorders. Neuroscience 440:337–359. https://doi.org/10.1016/j.neuroscience.2020.05.032
13. Smith RS, Walsh CA (2020) Ion channel functions in early brain development. Trends Neurosci 43:103–114. https://doi.org/10.1016/j.tins.2019.12.004
14. Gan J, Cai Q, Galler P (2019) Mapping the knowledge structure and trends of epilepsy genetics over the past decade A co-word analysis based on medical subject headings terms. Medicine (Baltimore) 98(32):e16782. https://doi.org/10.1097/MD.0000000000016782
15. D’Adamo MC, Liantonio A, Conte E et al (2020) Ion channels involved in neurodevelopmental disorders. Neuroscience 440:337–359. https://doi.org/10.1016/j.neuroscience.2020.05.032
16. Smith RS, Walsh CA (2020) Ion channel functions in early brain development. Trends Neurosci 43:103–114. https://doi.org/10.1016/j.tins.2019.12.004
17. Gan J, Cai Q, Galler P (2019) Mapping the knowledge structure and trends of epilepsy genetics over the past decade A co-word analysis based on medical subject headings terms. Medicine (Baltimore) 98(32):e16782. https://doi.org/10.1097/MD.0000000000016782
Berkovic FS, Petrou S (2014) KCNT1 gain of function in 2 epilepsy phenotypes is reversed by quinidine. Ann Neurol 75(4):581–590. https://doi.org/10.1002/ana.24128

27. Harkin LA, McMahon JM, Iona X, Dibbens L (2007) The spectrum of SCN1A-related infantile-epileptic encephalopathies. Brain 130(3):843–852. https://doi.org/10.1093/brain/awm002

28. Richards S, Aziz N, Bale S, Bik D, Das S, Gastier-Foster J, Grody WW, Hegde M, Lyon E, Spector E, Voelkerding K, Rehm LH (2015) Standards and guidelines for the interpretation of sequence variants: a joint consensus recommendation of the American College of Medical Genetics and Genomics and the Association for Molecular Pathology. Genet Med 17(5):405–424. https://doi.org/10.1038/gim.2015.30

29. Claes L, Del-Favero J, Ceulemans B, Lagae L, Van Broeckhoven C, De Jonghe P (2001) De novo mutations in the sodium-channel gene SCN1A cause severe myoclonic epilepsy of infancy. Am J Hum Genet 68:1327–1332. https://doi.org/10.1086/320609

30. Escayg A, Heils A, MacDonald BT, Haug K, Sander T, Meisler MH (2001) A novel SCN1A mutation associated with generalized epilepsy with febrile seizures plus—and prevalence of variants in patients with epilepsy. Am J Hum Genet 68:866–873. https://doi.org/10.1086/319524

31. Fujisawa T (2006) Clinical spectrum of mutations in SCN1A gene: severe myoclonic epilepsy in infancy and related epilepsies. Epilepsy Res 70(Suppl 1):S223–S230. https://doi.org/10.1016/j.eplepsyres.2006.01.019

32. Gennaro E, Santorelli FM, Bertini E, Buti D, Gaggero R, Gobbi G, Lini M, Granata T, Freire E, Parmeggiani A, Satriano P, Veggio P, Caradonna S, Bracaglia FD, Minetti C, Zara F (2006) Somatic and germine mosaicism in severe myoclonic epilepsy of infancy. Biochem Biophys Res Commun 341:489–493. https://doi.org/10.1016/j.bbrc.2005.12.209

33. Zuberi SM, Brunklau AJ, Birch R, Reavey E, Duncan J, Forbes GH (2011) Genotype-phenotype associations in SCN1A-related epilepsies. Neurology 76:934–941. https://doi.org/10.1212/WNL.0b013e318232309b

34. Martin HC, Kim GE, Pagnamenta AT (2014) Clinical whole-genome sequencing in severe early-onset epilepsy reveals new genes and improves molecular diagnosis. Hum Mol Genet 23(12):3200–3211. https://doi.org/10.1093/hmg/ddu380

35. Ishii A, Shioda M, Okamura A, Kidozora H, Sakauchi M, Shimada S, Shimizu T, Osawa T, Ikeda S, Yamashita Y, Yamamoto T (2013) A novel KCNT1 mutation in two sporadic cases with malignant migrating partial seizures in infancy. Genes 3(1):467–471. https://doi.org/10.3390/genes30100467

36. Lim CX, Ricos MG, Dibbens LM, Heron SE (2016) KCNT1 mutations in seizure disorders: the phenotypic spectrum and functional effects. J Med Genet 53(4):217–225. https://doi.org/10.1136/jmedgenet-2015-103508

37. Ohtba C, Kato M, Takahashi N et al (2015) De novo KCNT1 mutations in early-onset epileptic encephalopathy. Epilepsia 56(9):114–120. https://doi.org/10.1111/epi.13071

38. Moller RS, Heron SE, Larsen LHG (2015) Mutations in KCNT1 cause a spectrum of focal epilepsies. Epilepsia 56(9):114–120. https://doi.org/10.1111/epi.13071

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