Two siblings with early repolarization syndrome: clinical and genetic characterization by whole-exome sequencing

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Aims The early repolarization syndrome (ERS) can cause ventricular fibrillation (VF) and sudden death in young, otherwise healthy individuals. There are limited data suggesting that ERS might be heritable. The aim of this study was to characterize the clinical phenotype and to identify a causal variant in an affected family using an exome-sequencing approach.

Methods and results Early repolarization syndrome was diagnosed according to the recently proposed Shanghai ERS Score. After sequencing of known ERS candidate genes, whole-exome sequencing (WES) was performed. The index patient (23 years, female) showed a dynamic inferolateral early repolarization (ER) pattern and electrical storm with intracardiac VF. Isoproterenol enabled successful termination of electrical storm with no recurrence on hydroquinidine therapy during 33 months of follow-up. The index patient’s brother (25 years) had a persistent inferior ER pattern with malignant features and a history of syncope. Both parents were asymptomatic and showed no ER pattern. While there was no pathogenic variant in candidate genes, WES detected a novel missense variant affecting a highly conserved residue (p. H2245R) in the ANK3 gene encoding Ankyrin-G in the two siblings and the father.

Conclusion We identified two siblings with a malignant ERS phenotype sharing a novel ANK3 variant. A potentially pathogenic role of the novel ANK3 variant is suggested by the direct interaction of Ankyrin-G with the cardiac sodium channel, however, more patients with ANK3 variants and ERS would be required to establish ANK3 as novel ERS susceptibility gene. Our study provides additional evidence that ERS might be a heritable condition.

Keywords Early repolarization • Ventricular fibrillation • Electrical storm • Isoproterenol • Hydroquinidine • Exome sequencing • ANK3 • Ankyrin-G • Sodium channel
What's new?

- The early repolarization syndrome (ERS) is an extremely rare and malignant primary arrhythmia syndrome. Early repolarization syndrome might be heritable, however, there are no co-segregation data of genetic variants and ERS in affected families.
- We identified two siblings with symptomatic ERS and tried to identify a causal variant in the affected family by using a whole-exome sequencing (WES) approach.
- Whole-exome sequencing detected a novel, ultra-rare missense variant affecting a highly conserved residue in the ANK3 gene encoding Ankyrin-G in three out of four family members.
- There are clinical data supporting delayed depolarization as underlying mechanism in ERS and a potentially pathogenic role of the novel ANK3 variant in ERS is suggested by the direct interaction of Ankyrin-G with the cardiac sodium channel.

Introduction

The early repolarization (ER) pattern has been defined as J-point elevation of ≥0.1 mV in two continuous inferolateral leads on a standard 12-lead electrocardiogram (ECG) with either a distinct or partly buried J-wave. The malignant ER syndrome (ERS) is occurring in patients with an ER pattern that experience syncope, ventricular fibrillation (VF), or sudden cardiac death (SCD). A Shanghai ERS score of >5 points was required for a diagnosis of definitive ERS.1 A proband, his affected brother and the father as we did not have enough high-quality DNA of the mother for WES. The coding exons were captured using the Agilent SureSelect Human All Exon v5.1. The captured fragments were sequenced on the HiSeq2000 sequencer (Illumina San Diego, CA, USA) to an average depth of >50 reads per target base. The sequence reads were aligned to the human reference genome (UCSC NCBI37.1/hg19) using BWA and annotated by GATK for break-point identification. Variants was performed using SOAPsnp software for the single nucleotide variation. For insertion/deletion detection, sequence reads were aligned by BWA and annotated by GATK for break-point identification. Variants were annotated using ANNOVAR. Data analysis was performed in the statistical programming environment R version 3.2.1. Synonymous variations not located at splice sites and variants with a frequency >0.1% in the following publically available databases were excluded from further analysis: (i) the ExAC Browser, (ii) the National Heart, Lung, and Blood Institute (NHLBI) Exome Sequencing Project (ESP), (iii) Phase 1 version 3 of the 1000 Genomes project (data release October 2012), (iv) the 69 genomes from Complete Genomics, and (v) the Genome Aggregation Database (gnomAD). Sequential analysis steps focused on (i) Candidate gene analysis: known genes associated with ERS- and BrS (BrS, cardiac arrhythmias and cardiomyopathies, see Supplementary material online, S3 for the gene lists); (ii) recessive inheritance: shared homozygous variants in both siblings, heterozygous in the father and the mother; (iii) reduced penetrance: novel variants (i.e. ultra-rare variants shared by both affected siblings and inherited from an unaffected parent but not found in any of the databases).

Methods

The study was approved by the Institutional Ethics Committee on Human Research at the authors’ institution. All patients have given written informed consent for study participation, blood sampling, isolation of DNA, molecular genetic testing, and provision of their clinical data.

Clinical data analysis

Clinical characterization included medical history, analysis of 12-lead-, Holter-, and monitor ECGs, transthoracic echocardiography, coronary angiography, Ajmaline challenge (when deemed appropriate), exercise stress test, and Valsalva manoeuvre. The peak of an end-QRS notch and/or onset of an end-QRS slur, J peak (jp) was measured in standard 12-lead ECGs.1 A Shanghai ERS score of ≥5 points was required for a diagnosis of definitive ERS.10

Candidate gene analysis

DNA was extracted from peripheral blood lymphocytes of the two siblings and their parents and analysed by next-generation sequencing (NGS) using a customized (SureDesign) solution-capture enrichment strategy (Agilent SureSelect) on a MiSeqTM- or a NextSeqTM 500-System (Illumina, San Diego, CA, USA). Targeted sequencing of ERS- and Brugada syndrome (BrS)-related genes (GPD1L, SCN1B, KCNE3, SCN3B, KCNH2, RANGRF, KCNE1L, KOND3, HCN4, SLMP, TRPM4, SCN2B, SCN10A, KCNJ8, CACNA1C, CACNB2, CACNA2D1, SCN5A, and ABC9) was performed in the index patient. Target regions included coding exons (padding: ±25 bp). Sequencing was carried out on a MiSeqTM- or a NextSeqTM 500-System (Illumina, San Diego, CA, USA). Bioinformatic analyses were performed using BLAST+. Local analysis was performed using SOAPsnp software for the single nucleotide variation. For insertion/deletion detection, sequence reads were aligned by BWA and annotated by GATK for break-point identification. Variants were annotated using ANNOVAR. Data analysis was performed in the statistical programming environment R version 3.2.1. Synonymous variations not located at splice sites and variants with a frequency >0.1% in the following publically available databases were excluded from further analysis: (i) the ExAC Browser, (ii) the National Heart, Lung, and Blood Institute (NHLBI) Exome Sequencing Project (ESP), (iii) Phase 1 version 3 of the 1000 Genomes project (data release October 2012), (iv) the 69 genomes from Complete Genomics, and (v) the Genome Aggregation Database (gnomAD). Sequential analysis steps focused on (i) Candidate gene analysis: known genes associated with ERS- and BrS (BrS, cardiac arrhythmias and cardiomyopathies, see Supplementary material online, S3 for the gene lists); (ii) recessive inheritance: shared homozygous variants in both siblings, heterozygous in the father and the mother; (iii) reduced penetrance: novel variants (i.e. ultra-rare variants shared by both affected siblings and inherited from an unaffected parent but not found in any of the databases).

Results

Clinical phenotype

The index patient, individual II-2 in the pedigree (Figure 1A), experienced out-of-hospital-cardiac arrest at age 23 years in the early morning at home while sleeping. Basic life support was provided by her mother. Upon arrival of the emergency physician, the initially recorded rhythm was VF. After transfer to University Hospital Freiburg ischaemic and structural heart disease were ruled out by transthoracic echocardiography and coronary angiography. Due to recurrent VF, the patient underwent early percutaneous venoarterial extracorporeal membrane oxygenation (VA-ECMO) implantation.

Whole-exome sequencing

Whole-exome sequencing was initially carried out on the proband, her affected brother and the father as we did not have enough high-quality DNA of the mother for WES. The coding exons were captured using the Agilent SureSelect Human All Exon v5.1. The captured fragments were sequenced on the HiSeq2000 sequencer (Illumina San Diego, CA, USA) to an average depth of >50 reads per target base. The sequence reads were aligned to the human reference genome (UCSC NCBI37.1/hg19) using SOAPsnp software for the single nucleotide variation. For insertion/deletion detection, sequence reads were aligned by BWA and annotated by GATK for break-point identification. Variants were annotated using ANNOVAR. Data analysis was performed in the statistical programming environment R version 3.2.1. Synonymous variations not located at splice sites and variants with a frequency >0.1% in the following publically available databases were excluded from further analysis: (i) the ExAC Browser, (ii) the National Heart, Lung, and Blood Institute (NHLBI) Exome Sequencing Project (ESP), (iii) Phase 1 version 3 of the 1000 Genomes project (data release October 2012), (iv) the 69 genomes from Complete Genomics, and (v) the Genome Aggregation Database (gnomAD). Sequential analysis steps focused on (i) Candidate gene analysis: known genes associated with ERS- and BrS (BrS, cardiac arrhythmias and cardiomyopathies, see Supplementary material online, S3 for the gene lists); (ii) recessive inheritance: shared homozygous variants in both siblings, heterozygous in the father and the mother; (iii) reduced penetrance: novel variants (i.e. ultra-rare variants shared by both affected siblings and inherited from an unaffected parent but not found in any of the databases).
The initial 12-lead ECG showed normal sinus rhythm with a QTc of 421 ms, and an inferolateral ER pattern with Jp amplitudes $\geq 0.1$ mV (Figure 1C). The telemetry monitor revealed a bradycardia-dependent inferior ER pattern with frequent premature ventricular complexes triggering polymorphic ventricular tachycardia (pVT) and recurrent VF (Figure 2). Conventional anti-arrhythmic drugs failed and the patient was eventually rescued from intractable VF with isoproterenol which enabled successful defibrillation and suppression of ER and VF—after more than 1 h of ongoing VF (Figure 3). During isoproterenol weaning, monomorphic couplets recurred and isoproterenol was replaced with hydroquinidine 300 mg bid, which again suppressed couplets and ER (Supplementary material online, S1). The patient recovered completely without any sequelae and was implanted with a dual-chamber implantable cardioverter-defibrillator (ICD) with atrial pacing set at 70 b.p.m. There has been no VF recurrence on hydroquinidine therapy during 33 months of follow-up. Ajmaline challenge including high precordial leads placed in 3rd and 2nd intercostal space in (para-)sternal positions was negative. The Shanghai ERS score of the index patient was consistent with a definitive diagnosis of ERS (cardiac arrest/documented VF: 3 points; ER $\geq 0.2$ mV in $\geq 2$ lateral ECG leads with horizontal ST-segment: 2 points).

The index patient’s brother (25 years) demonstrated a persistent inferior ER pattern with high inferior Jp amplitudes $\geq 0.35$ mV and horizontal ST-segments (Figure 1B). Ajmaline challenge with precordial leads placed as in the index was negative. The past medical history was significant for a witnessed arrhythmic syncope at rest without any prodrome or trigger at age 14 years. The Shanghai ERS score was calculated to be 6 points (suspected arrhythmic syncope: 2 points; ER $\geq 0.2$ mV in $\geq 2$ inferior ECG leads with horizontal ST-segment: 2 points; relative with definitive ERS: 2 points) consistent with a definitive diagnosis of ERS. In addition, the ER pattern showed a dynamic augmentation during Valsalva (Supplementary material online, S2). Based on a malignant ER pattern, a strong family history of ERS, and a history of arrhythmic syncope at rest the patient was deemed at increased risk of pVT/VF and a subcutaneous ICD was implanted. The patient has not received any shocks during follow-up.

Both parents were asymptomatic, they had a negative (extended) family history for unexplained syncope or SCD and showed no ER pattern on several 12-lead ECGs at baseline and during Valsalva.
We did not perform Ajmaline challenges in the parents due to the negative Ajmaline test results in both siblings.

**Genetic analysis**

A diagnostic screening by NGS showed no potentially causal variant in GPD1L, SCN1B, KCNE3, KCN2B, RANGRF, KCNE1L, KCND3, HCN4, SLMAP, TRPM4, SCN2B, SCN10A, KCNJ8, CACNA1C, CACNB2, CACNA2D1, SCN5A, and ABCC9 in the proband. Whole-exome sequencing was performed in both affected siblings and the unaffected father. The analysis strategy for the WES data consisted of three steps: (i) analysis of candidate gene panels, (ii) analysis for potential recessive inheritance; (iii) analysis of potentially dominant inheritance with reduced penetrance. No rare variants shared by both affected siblings were identified in genes associated with ERS in the literature, nor in genes associated with cardiac arrhythmias or cardiomyopathy (Supplementary material online, S3). Both siblings shared 1 rare homozygous variant in the gene FADS6 which was shared homozygously by the unaffected father as well and therefore did not fit a putative recessive pattern of inheritance with complete penetrance. No compound heterozygous variants were identified. We next investigated whether any ultra-rare variants (not present in all reference databases) were shared by both siblings. Thirteen variants fitted these criteria (Supplementary material online, S3). Among these was a novel variant in the gene ANK3 (NM_020987;...
exon 37; c. A6734G; p. H2245R), shared by both siblings and inherited from the unaffected father which was absent in the mother. Histidine 2245 of ANK3 is located in a highly conserved domain of the protein, it is unchanged from Xenopus to fish, birds, and mammals (Figure 4) and the CADD-phred score of this variant is 14.77 indicating a strong conservation and large predicted effect on the protein. None of the other variants were in genes expressed in the heart (GTEX tpm >5), with a CADD-phred score >10, in genes with a significant constraint based on gnomAD pLoF (<0.05).

**Discussion**

In this study, we made an attempt to investigate the genetic background of ERS in a small affected family by candidate gene analysis and a WES approach. The clinical presentation and dynamic ER pattern—aggravated by pauses, bradycardia, and changes in vagal tone (Valsalva) and resolved with sympathetic activation by isoproterenol—support the diagnosis of ERS in the two siblings, which is reflected by high ERS scores. Thus, our study provides additional evidence that ERS could be a heritable condition.

The underlying mechanism of ER is a matter of debate and currently, it is not clear whether ERS involves disturbed depolarization, repolarization, or both.11,12 The results of genetic and functional studies and pharmacological interventions with isoproterenol and hydroquinidine are compatible with the cellular model of ER, i.e., an imbalance between depolarizing currents (sodium and L-type calcium currents, \(I_{Na}\) and \(I_{Ca,L}\)) and repolarizing transient outward potassium current (\(I_{to}\)) during the early phase of the epicardial action potential. However, the magnitude of these currents and the resulting action potential morphology (i.e., the voltage-time integral ‘area under the curve’) also determine the safety of conduction and hence conduction velocity.13 As isoproterenol and hydroquinidine increase the area under the curve, such pharmacological interventions will also improve conduction which is compatible with disturbed depolarization in ERS similar to BrS.13,14 Interestingly, the index patient’s brother showed a fragmented QRS in precordial and lateral leads compatible with abnormal depolarization (Supplementary material online, S4).

While we did not find any pathogenic variant in ERS candidate genes, WES detected a novel ultra-rare missense variant affecting a highly conserved residue (p. H2245R) in the ANK3 gene encoding Ankyrin-G in the two symptomatic siblings and the unaffected father. A potentially pathogenic role of the novel ANK3 variant is suggested by the direct interaction of Ankyrin-G with the cardiac sodium channel: ANK3 encodes Ankyrin-G which is required for targeting Nav1.5 to the intercalated disc.15–17 In animals, ankyrin-G expression is required for the targeting of Nav1.5 and CaMKII to the intercalated disc16 and Nav1.5 expression and \(I_{Na}\) are reduced in myocytes derived from mice with selective loss of cardiomyocyte ankyrin-G (cKO).16 Priori et al identified an individual with BrS harbouring an SCN5A variant in the Nav1.5 motif that interacts with ankyrin-G. In vitro binding assays show that the p. E1053K variant disrupts the interaction between Nav1.5 and ankyrin-G resulting in abnormal targeting in myocytes.18 Interestingly, Ankyrin variants have also been found in two young, unrelated individuals with VF storm due to possible ERS, one of them with a family history of SCD.19 In summary, as Ankyrins are a critical part of the sodium channel complex, the novel ANK3 variant could impair conduction via loss-of-sodium channel function20 which may contribute to disturbed depolarization in ERS. However, more patients with variants in ANK3 (and familial) ERS would be required to establish ANK3 as a novel ERS susceptibility gene. In addition, the presence of the novel ANK3 variant in the unaffected father may support a more complex genetic architecture of ERS (e.g., additional genetic variants/modifiers), may be due to the fact that the novel variant is not disease causing but may also point to incomplete disease penetrance or variable disease expression which has been described in families with ERS.6

**Limitations**

The small number of affected individuals in our study does not allow to conclude a causal role of the novel variant in this family and in ERS in general. Functional studies, e.g. iPS cells, would be required to assess the effect on ionic currents, e.g. a reduction in \(I_{Na}\), and could be of additional value to support the pathogenicity of the novel ANK3 variant.
Conclusion

Our study provides additional evidence that ERS could be a heritable condition with variable penetrance. This study is, however, limited by the small number of affected individuals which does not allow a conclusion regarding a pathogenic role of the novel ANK3 variant in this family and in ERS in general. Additional patients with variants in ANK3 and ERS as well as functional studies are required to establish a disease-modifying or pathogenic role of ANK3 in ERS.

Supplementary material

Supplementary material is available at Europace online.

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Conflict of interest: none declared.

Data availability

The data underlying this article will be shared on reasonable request to the corresponding authors.

References