Inherited conditions of arrhythmia: translating disease mechanisms to patient management

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Inherited arrhythmias may be caused by mutations in genes encoding cardiac ion channels and associated proteins (primary electrical disorders), and may occur in the setting of hereditary cardiomyopathies secondary to mutations in cardiac structural proteins. Together, they may contribute to up to 15–20% of all sudden cardiac deaths (SCDs). Progress in genetic, molecular, and (electro)physiological aspects of inherited arrhythmia disorders has enabled the identification of diagnostic and therapeutic strategies. However, insight into disease mechanisms and arrhythmia triggers is still limited, precluding the development of mechanism-driven therapies. In addition, inherited arrhythmia syndromes are often associated with reduced penetrance and variability in disease expressivity and severity. Hence, predicting who is most at risk for SCD remains difficult. In this Spotlight Issue, a series of reviews highlight the current state of the art for the various inherited arrhythmia disorders, review novel targets for risk stratification and therapy, and describe the remaining challenges and future perspectives.

In line with their role in action potential formation, mutations in ion channel genes are well-established causes of inherited arrhythmia syndromes associated with altered depolarization and/or repolarization. A wide range of molecularly diverse potassium channels are present in the myocardium, which together maintain the resting membrane potential and mediate action potential repolarization. Mutations in genes encoding these channels can impair their assembly, trafficking, and gating, leading to long QT syndrome types 1, 2, and 3 (LQT1, LQT2, LQT3), Brugada syndrome (BrS), cardiac conduction disease, atrial fibrillation, and sick sinus syndrome. In their review, Rivaud et al. discuss the multifunctionality of sodium channels and propose a novel classification of NaV1.5 (dys)function. Reduced peak sodium current or increased late sodium current (‘direct ionic’ effects) consequent to SCN5A mutations are well-established causes of pro-arrhythmic conduction slowing and repolarization disturbances, respectively. However, cardiac abnormalities are increasingly reported in SCN5A mutation carriers, including cardiac fibrosis, dilated cardiomyopathy, and arrhythmogenic (right ventricular) cardiomyopathy (ACM/ARVC). Such structural remodelling may occur consequent to alterations in intracellular sodium and calcium homeostasis (‘indirect ionic’ effects), or secondary to disrupted interactions of NaV1.5 with partner proteins within the macromolecular complex (‘non-ionic’ effects). Since these non-canonical actions of NaV1.5 may contribute significantly to arrhythmogenesis, they clearly warrant further exploration.

In addition to ion channelopathies, arrhythmias and SCD also occur in the setting of hereditary cardiomyopathies, including hypertrophic cardiomyopathy (HCM), dilated cardiomyopathy (DCM), arrhythmogenic cardiomyopathy (ACM), and left ventricular non-compaction cardiomyopathy (LVNC). As reviewed by Marian et al., these cardiomyopathies are predominantly caused by mutations in genes encoding sarcomeric proteins (HCM), cytoskeletal proteins (DCM), and desmosomal proteins (ACM). Pro-arrhythmic mechanisms include cardiac hypertrophy and dilatation, myocardial fibrosis, ion channel and connexin remodelling, and intracellular calcium dysregulation. Interestingly, mutations in a number of genes encoding ion channels or transporters, including SCN5A, KCNJ2, PLN (phospholamban), and RYR2 (ryanodine receptor 2) have been associated with DCM, ACM, and/or LVNC. Conversely, mutations in the LMNA gene encoding lamin A/C lead to cardiac conduction defects and modulation of ion currents, or rescue of trafficking-deficient channels.

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ventricular arrhythmias, and mutations in titin (TTN) to atrial fibrillation. Hence, inherited cardiomyopathies and cardiac arrhythmias are closely interrelated, and may have in part a shared genetic aetiology. This is further exemplified by ACM/ARVC, which is predominantly caused by mutations in desmosomal genes but which was recently also linked to mutations in SCN5A. The pathogenesis of ACM includes intercalated disc remodelling, cardiomyocyte loss, adiposis, and inflammation. As discussed by Van der Voorn et al.,7 novel anti-fibrotic and anti-inflammatory strategies are currently being investigated in available mouse, zebrafish, and hiPSC-CM ACM disease models. Various mechanisms may contribute to arrhythmias in ACM: cardiac structural alterations and fibrosis formation may set the stage for re-entrant based arrhythmias in advanced disease stages, whereas Na\textsubscript{v}1.5 remodelling and calcium dysregulation may occur in early disease stages and cause arrhythmias prior to overt cardiomyopathic changes. Novel strategies enabling early detection of pathological remodelling and hence risk stratification in ACM patients include (bio)markers of fibrosis and inflammation, desmosomal protein remodelling in buccal mucosa smears, circulating desmoglein-2 autoantibodies, and assessment of cardiac conduction delay through high resolution imaging and sodium channel blocker challenge. Abnormalities in sodium current and calcium homeostasis also contribute to arrhythmogenesis in the setting of HCM, as reviewed by Coppini et al.8 HCM is a major cause of SCD in young individuals (occurring during exercise or at rest), but the factors predisposing to ventricular arrhythmias remain incompletely known and hence risk stratification options are limited. While structural alterations may provide a pro-arrhythmic substrate, a clear association between e.g. cardiac fibrosis and arrhythmia risk is lacking in HCM patients. More recently, the use of mouse, rabbit, and hiPSC-CM models of HCM-related mutations have provided essential mechanistic insight into electrophysiological remodelling on the cardiomyocyte level. Decreased potassium currents, increased L-type calcium current and enhanced late sodium current have been shown to underlie action potential prolongation and EAD formation, attributed at least in part to a sustained CamKII activation. In addition, global dysregulation of intracellular calcium homeostasis occurs in HCM cardiomyocytes, secondary to alterations in L-type calcium current and sarcoplasmic reticulum and NCX function. Diastolic calcium levels are furthermore elevated consequent to altered sarcomere function and set the stage for DAD formation and triggered activity. Pharmacological late sodium current inhibition has been shown to improve calcium handling and reduce pro-arrhythmia in HCM cardiomyocytes, indicating a potential therapeutic strategy.

While genetic studies in the inherited arrhythmia field have traditionally focused on Mendelian disorders, it has become increasingly clear that the genetic basis for these disorders is often far more complex.1 As discussed above, mutations in one gene can lead to multiple disease phenotypes, and conversely one disease can be caused by mutations in multiple genes. In addition, genetic modifiers likely contribute to the reduced penetrance and variable disease severity observed in patients. In some disorders, such as for instance Brugada syndrome (BrS), an oligogenic or polygenic basis has been demonstrated, with multiple common or rare genetic variants in aggregate predisposing to the disease. In their review, Glinge et al. discuss the use of genome-wide association studies (GWAS) for the identification of genetic variants (single nucleotide polymorphisms) that govern interindividual variability in electrocardiographic parameters in the general population.9 Identified variants can be used to explore their possible role in modifying disease susceptibility and

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**Inherited Cardiomyopathies and Primary Electrical Disorders**

**State of the Art**
- Monogenic inheritance
- Limited therapeutic options
- Models: transgenic animals, hiPSC-CMs
- Genetic overlap:

**Challenges**
- Novel disease mechanisms
- Disease modifiers
- Complex genetics (oligo-/polygenic)
- Functional validation of variants
- Arrhythmia triggers
- SCD risk prediction
- Mechanism-driven therapy
- Gene therapy

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![Figure 1](image.png) Inherited cardiomyopathies and primary electrical disorders: state of the art and challenges.
efforts will no doubt continue to facilitate development of improved strategies for diagnosis, risk stratification, prevention, and treatment of patients with inherited arrhythmias.

Conflict of interest: none declared.

Funding
This work was supported by a grant from the Netherlands Cardio Vascular Research Initiative (CVON): the Dutch Heart Foundation, Dutch Federation of University Medical Centres, the Netherlands Organization for Health Research and Development and the Royal Netherlands Academy of Sciences (CVON-eDETECT 2015-12, CVON-PREDICT2 2018-30) to C.A.R.

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