The pacemaker cells of the cardiac sinoatrial node (SAN) are essential for normal cardiac automaticity. Dysfunction in cardiac pacemaking results in human sinoatrial node dysfunction (SND). SND more generally occurs in the elderly population and is associated with impaired pacemaker function causing abnormal heart rhythm. Individuals with SND have a variety of symptoms including sinus bradycardia, sinus arrest, SAN block, bradycardia/tachycardia syndrome, and syncope. Importantly, individuals with SND report chronotropic incompetence in response to stress and/or exercise. SND may be genetic or secondary to systemic or cardiovascular conditions. Current management of patients with SND is limited to the relief of arrhythmia symptoms and pacemaker implantation if indicated. Lack of effective therapeutic measures that target the underlying causes of SND renders management of these patients challenging due to its progressive nature and has highlighted a critical need to improve our understanding of its underlying mechanistic basis of SND. This review focuses on current information on the genetics underlying SND, followed by future implications of this knowledge in the management of individuals with SND.

Keywords: genetics, sick sinus syndrome, sinoatrial node dysfunction, atrial fibrillation, calsequestrin-2, GIRK4, HCN4, Na\textsubscript{v}1.5

INTRODUCTION

Automaticity is the primary function of the sinoatrial node (SAN). SAN pacemaker cells have the shortest depolarization phase of the action potential and the fastest firing rate, making the SAN the dominant pacemaker of the heart (Kennedy et al., 2016). The SAN initiates an electrical impulse that propagates throughout the heart, establishing a normal heart rhythm. The sinoatrial node is surrounded by a collagen frame, which helps protect SAN cells against ectopic activation from other atrial cells (James, 1977; Kalyanasundaram et al., 2019). The SAN location was initially reported
near the epicardium between the superior vena cava and the right atrium in the posterior right atrial wall (Keith and Flack, 1907). The human SAN was later described to commonly exist laterally and inferior to the terminal crest. Infrequently, the SAN can exist at the junction between the superior caval vein and right atrium, or even across the terminal crest into the interatrial groove (Anderson et al., 1979; Sánchez-Quintana et al., 2005; Dobrzenski et al., 2007). Recently, evidence has suggested the simultaneous existence of superior and inferior SAN in the human heart located near the superior and inferior venae cavae, respectively. The superior SAN is suggested to generally control higher heart rate, while the inferior (“back-up”) SAN is suggested to control lower heart rate. Optical mapping data shows that the heart rate control distinctly jumps between cluster control, rather than a gradient transition, during physiologic heart rate changes (Brennan et al., 2020). The SAN primarily contains three specialized cell types: pale cells (P cells), transitional cells (T cells), and recently described “fibroblast-like” cells, intertwined in collagen, fibroblasts, fatty tissue, nerves, and capillaries (James, 1977; Balbi et al., 2011; Csepe et al., 2015). P cells are localized to the central region of the SAN and have a single membrane with very little membrane specialization resulting in poorly defined intercalated discs, few desmosomes, and few gap junctions (Ho and Sánchez-Quintana, 2016). Gap junctions are rare in P cells, and junctions are only formed between P cells or T cells. P cells do not form junctions with other atrial myocytes (James, 1977). Automaticity arises from P cells (James et al., 1966; Woods et al., 1976). Interestingly, transitional cells, located in the periphery of the SAN, act as the distributors of action potentials to atrial myocardium and internodal conduction pathways (Boyett et al., 2000). T cells are a loosely defined, varied group of cells. Many T cells look like P cells with reduced myofibrils, while others closely resemble atrial myocytes (James et al., 1966). Finally, the function of “fibroblast-like” cells has yet to be defined, but presumably, these cells play a role in maintaining the structure of the SAN (Balbi et al., 2011). The SAN was initially proposed to propagate action potentials to the atrioventricular (AV) node via three tracts: the anterior (which splits into two bundles), middle, and posterior internodal tracts (James, 1963; Kennedy et al., 2016). However, recent studies failed to confirm this hypothesis. Interestingly, current optical mapping studies using rat and human atrial tissues suggest the presence of more than a single definitive pacemaker site. Overall, the existence of clearly defined internodal tracts has not been confirmed via optical and electrical mapping (Boineau et al., 1988; Brennan et al., 2020).

The SAN receives blood flow through the sinus node artery, the largest atrial coronary branch, which originates from the left or right coronary artery (Verhaeghe and van der Hauwaert, 1967). Of note, parasympathetic modulation of the heart is primarily through the vagal postganglionic pathways along the sulcus terminalis in the subepicardial region adjacent to the SAN artery (Bluemel et al., 1990). Innervation is both parasympathetic and sympathetic to the SAN, yet neither innervation pathway has any direct contact with P or T cells (Balbi et al., 2011). The intrinsic cardiac nervous system is mainly composed of cardiac ganglia that play a significant role in the integration of cardiac electrophysiology (Fedele and Brand, 2020).

Interestingly, it has been suggested that the total amount of SAN cells is inversely proportional to the age of an individual, suggesting that the number of SAN cells and the volume of SAN cells decrease over time (Thery et al., 1977; Shiraishi et al., 1992). However, Alings et al. (1995) found that the SAN does not change in dimensions during an entire adult life span. Although relative collagen increases from childhood to adulthood, collagen levels do not change once adulthood is reached. Instead, the structure of the collagen changes throughout adulthood (Alings et al., 1995).

SND occurs in one of every 600 cardiac patients above the age of 65 (Rodriguez and Schocken, 1990). Symptomatic SND is the most common rationale for permanent pacemaker implantation (Kusumoto et al., 2019). In general, “sick sinus syndrome” (SSS), sinus node dysfunction (SND), and sinoatrial node dysfunction (also SND) are used interchangeably in the literature. SND is usually accompanied by structural abnormalities. Symptoms of SND include sinus bradycardia [heart rate less than 60 beats per minute (bpm)], sinus arrest (total absence of atrial or ventricular activity), SA block, and bradycardia/tachycardia syndrome (cycling between supraventricular tachyarrhythmias and sinus bradycardia) (Alpert and Flaker, 1983). Syncope has also been reported by about half of patients with SND (Menozzi et al., 1998; Briggle et al., 2013). Although structural defects accompany most forms of SND, syncope and bradycardia associated with SND have also been observed in patients with normal cardiac anatomy (Yabek et al., 1982). SND is generally associated with impaired pacemaker function causing abnormal heart rhythm, but SND can also be associated with impulse transmission issues from apoptosis in the AV node, sinus node, and internodal pathways, which may cause death from complete heart block (James et al., 1996). Further, blockage of the sinus node artery preventing blood flow to the SAN cells can result in SND and sudden death (Jing and Hu, 1997).

This review focuses on current information on the genetics underlying SND, followed by a brief overview of the recent update in clinical management of patients with SND. A better understanding of the genetics and the molecular mechanisms of the SAN and SND will improve current diagnostic measures and identify alternative therapeutic approaches. The genes currently implicated in human SND are summarized in Table 1. Although not the focus of this review, SND is also commonly a secondary symptom to other systemic and cardiovascular conditions. Congestive heart failure in particular results in SAN remodeling and a reduction in resting heart rate, which can result in SND (Sanders et al., 2004). Atrial fibrillation (AF) and atrial tachycardia, cardiac transplantation, drug toxicity, hyperinsulinemia and insulin resistance (diabetes type II), sinus node artery obstruction, hyperparathyroidism, intracranial conditions, epilepsy, myxedema coma, cardiac lymphoma, infections, and myocardial infarction are all potential procedures, diseases, or conditions that can result in secondary SND (Lown, 1967; Bexton et al., 1984; Bigger and Sahar, 1987; Wasada et al., 1995; Hasdemir et al., 2003; Shah et al., 2004; Dadlani et al., 2010; Ravindran et al., 2016; Kondo et al., 2020; Kousa et al., 2020;
Genetic Complexity of Sinoatrial Node Dysfunction

SND mainly affects the older population, although it can affect people at any age. Most cases of SND are not inherited. However, several genes coding for ion channels such as HCN4 and SCNA5A, cytoskeletal proteins, and proteins intricate to cardiac development have been linked to SND (de Ponti et al., 2018). Gene products associated with SND in the SAN are shown in Figure 1. Major genes linked to SND etiology, as well as human studies and animal models used to characterize the SND phenotype, will be discussed below. Mouse models are a common tool for in vivo studies of various cardiac proteins and genes. Although heart size and heart rate are significantly different between humans and mice, the human and murine hearts develop and function similarly (Wessels and Sedmera, 2003). Still, comparisons between the two species are not perfect. Note that many mouse models are full body or cardiac-specific knockouts of a gene, while human gene variants most commonly only have a single nucleotide change, which may still result in a fully transcribed protein. CRISPR/Cas9 technology, which allows the development of more humanized animal models via induction of specific point mutations, remains one of the greatest tools for improving translational animal studies going forward.

Genetic complexity and pleiotropy remain a big challenge to proper SND/SSS diagnosis with likely multiple complexities involved including genetic variation, underlying conditions, and environmental factors. With numerous SA-expressed gene variants associated with SND, and only moderate penetrance, SND can continue within families for generations before diagnosis (Abe et al., 2014). Further, compound variants for prominent SND-implicated genes seem to increase penetrance, emphasizing SND as an oligogenic disease (Baskar et al., 2014; de Filippo et al., 2015; Sacilotto et al., 2017).

| Protein | Gene | Associated cardiac diseases | References |
|---------|------|-----------------------------|------------|
| Calsequestrin-2 | CASQ2 | SND/bradycardia, catecholaminergic polymorphic ventricular tachycardia (CPVT), atrial arrhythmias | Reid et al., 1975; Leenhardt et al., 1995; Sumitomo et al., 2007; Weelke et al., 2014; Olekhov et al., 2015 |
| Ryanodine receptor 2 | RYR2 | SND/bradycardia, CPVT, atrial arrhythmias | Sumitomo et al., 2007; Domingo et al., 2015; Miyata et al., 2018; Dharmawan et al., 2019 |
| G protein-activated inward rectifier potassium channel 4 | KCNJ5 | SND/bradycardia, atrial arrhythmias, long QT syndrome type 13, Andersen–Tawil syndrome | Dobrev et al., 2001; Zhang et al., 2009; Yang et al., 2010; Wang et al., 2013; Kokunai et al., 2014; Kuß et al., 2019 |
| Guanine nucleotide-binding protein subunit beta-2/5 | GNBP2/GNBP5 | SND/bradycardia, cognitive disability, cardiac conduction abnormalities | Lodder et al., 2016; Stallmeyer et al., 2017; Fukuda et al., 2020 |
| Sodium/calcium exchanger 1 | SLC6A1 | Conduction disorders (PR and QT prolongation), ventricular arrhythmias, Kawasaki disease | Kim et al., 2012; Hong et al., 2014 |
| Sodium voltage-gated channel alpha subunit 5 | SCN5A | SND/bradycardia, long QT syndrome type 3, Brugada syndrome, dilated cardiomyopathy, conduction disorders, infant death syndrome | van den Berg et al., 2001; Benson et al., 2003; McNair et al., 2004; Smits et al., 2005; Nakajima et al., 2013; Freyermuth et al., 2016; Yang et al., 2017; Denti et al., 2018 |
| Hyperpolarization activated cyclic nucleotide gated potassium channel 4 | HCN4 | SND/bradycardia, ventricular arrhythmias, left ventricular non-compaction | Schulze-Bahr et al., 2003; Ueda et al., 2004; Nof et al., 2007; Milano et al., 2014 |
| Ankyrin-B | ANK2 | SND/bradycardia, CPVT, atrial arrhythmias, arrhythmogenic cardiomyopathy | Curran and Mohler, 2011; Roberts et al., 2019 |
| Myosin heavy chain 6 | MYH6 | SND/bradycardia, aorta coarctation, ventricular arrhythmias | Holm et al., 2011; Lam et al., 2015; Björnsson et al., 2016 |
| Lamin A | LMNA | SND/bradycardia, dilated cardiomyopathy, conduction disorders | MacLeod et al., 2003; Zaragoza et al., 2016; Lin et al., 2018; Yokokawa et al., 2019 |
| L-type calcium channel subunit Ca2.1.3 | CACNA1D | Sinoatrial node dysfunction and deafness (SANDO) | Baig et al., 2011; Liagat et al., 2019 |
| Short-stature homeobox 2 | SHOX2 | SND/bradycardia, atrial arrhythmias | Hoffmann et al., 2016, 2019; Li et al., 2016 |
Calsequestrin-2 (CASQ2) and Ryanodine Receptor 2 (RYR2)

Background
Cardiac calsequestrin, calsequestrin-2 (Casq2), localized to the sarcoplasmic reticulum (SR), is a low-affinity, high-capacity, Ca\(^{2+}\)-binding protein involved in the ability of the SR to preserve and release Ca\(^{2+}\) in cardiac myocytes (Murphy et al., 2011). Voltage-gated Ca\(^{2+}\) channels (L-type calcium channels) create an initial Ca\(^{2+}\) influx into the cell, which causes RyR2 receptors to release more calcium from the SR in a process called calcium-induced calcium release. Calsequestrin, the major Ca\(^{2+}\)-binding protein in the SR, is anchored to the membrane of the SR by RyR2 (either directly or indirectly via triadin and junctin) and regulates the release of Ca\(^{2+}\) through the RyR2 channel (Beard et al., 2004). Importantly, calsequestrin, the ryanodine receptor, junctin, and triadin are all necessary for proper calcium cycling (Zhang et al., 1997).

Human Studies
Loss-of-function variants in CASQ2 can result in catecholaminergic polymorphic ventricular tachycardia (CPVT), bradycardia, and atrial arrhythmias (Glukhov et al., 2015). CPVT was first identified in 1975 as a case report of “bidirectional tachycardia” induced by physical effort and stress in a 6-year-old female with no structural abnormalities (Reid et al., 1975). A later follow-up of 21 children over 7 years properly emphasized CPVT as a separate entity of ventricular tachycardia in children, emphasizing the dangers of misdiagnosis with the lurking potential for spontaneous sudden cardiac death (SCD) without immediate therapy. Leenhardt et al. (1995) showed the adrenergic-induced ventricular arrhythmias to be preventable with proper β-blocker treatment.

The CPVT phenotype from CASQ2 variants is attributed to an increase in expression of calreticulin and RyR2 (Song et al., 2007). CPVT is also commonly induced by RYR2 variants and has been highly associated with bradycardia (Miyata et al., 2018). An exon 3 deletion of RYR2 (c.169-353_273 + 657del) was found in a family with a history of SND and CPVT (Dharmawan et al., 2019). Exon 3 deletions were identified in two additional families that showed symptoms of SND (Bhuiyan et al., 2007). Further, a RyR2 (R420Q) missense mutation was associated with sinus bradycardia and atrial arrhythmias (Domingo et al., 2015). Although CPVT is well documented across CASQ2 and RYR2 variants, SAN dysfunction and atrial arrhythmias have been identified in CPVT patients, signifying the importance of calsequestrin-2 and ryanodine receptor 2 in proper SAN functioning (Sumitomo et al., 2007). Additionally, a study across
303 patients with lone AF found rare CASQ2 variants, implicating CASQ2 in AF as well (Weeke et al., 2014).

**Mouse Models**

Glukhov et al. (2015) described the first Casq2-null (Casq2<sup>−/−</sup>) mouse model with SAN defects. The phenotype included bradycardia, RR interval variability, SAN conduction abnormalities, and abnormally high atrial ectopic activity resulting in AF, implicating calsequestrin-2 in SAN functioning. Increased fibrosis in the pacemaking complex was observed in Casq2-deficient mice. At the cellular level, lack of functioning Casq2 caused abnormal calcium release from the SR and increased diastolic calcium concentration, resulting in a delay between action potential and transient calcium upstrokes (Glukhov et al., 2015). This disruption in intracellular calcium cycling explains the SAN, atrial myocyte, and ventricular myocyte dysfunction seen in individuals with CASQ2 loss-of-function variants.

Atrial burst pacing induced atrial flutter and AF in Casq2<sup>−/−</sup> mice versus wild-type (WT) mice, and isolated Casq2-deficient hearts showed ectopic foci from the pulmonary vein region when visualized with atrial optical voltage maps. Isolated hearts from Casq2-null mice also experienced diastolic subthreshold spontaneous Ca elevations (SCaEs) and delayed afterdepolarizations (DADs) when AF was not reached, but R-propafenone (RyR2 and Na<sup>+</sup> channel blocker) was shown to prevent AF, DADs, and SCaEs in these mice. The authors attributed this success mostly to the inhibition of RyR2, as an equipotent Na<sub>c</sub> channel inhibitor did not reach the same result (Faggioni et al., 2014). Notably, atrial overdrive pacing suppressed ventricular arrhythmias in a Casq2-null mouse model and therefore could provide a new therapeutic option for CPVT (Faggioni et al., 2013). Interestingly, a RyR2 R4496C homozygous mouse showed sinus pausing, atrial arrhythmias, and reduced SAN automaticity. This variant is associated with enhanced RyR2 activity and supports a link between RyR2 variants in humans with SND (Zhang et al., 2013). In general, loss-of-function Casq2 variants and gain-of-function RyR2 variants both contribute to unreliable and irregular calcium cycling in the SAN, atria, and ventricles. This likely translates to human variants and provides some insight into the cause of CPVT, AF, and SND seen in carriers of CASQ2 and RyR2 variants.

**G Protein-Activated Inward Rectifier Potassium Channel 4 (KCNJ5)**

**Background**

Activation of G protein via a transmitter is a common type of cell–cell communication. In general, a neurotransmitter binds to a seven transmembrane receptor on the outside of the cell, which results in the exchange of GDP for GTP on the inward side of the receptor, allowing the dissociation of the heterotrimeric G-protein subunits, which then further act as effectors (Gilman, 1987; Neer, 1995; Mark and Herlitze, 2000). For heart rate modulation, the G<sub>βγ</sub> (as opposed to the Gα) subunit specifically activates the K<sub>ach</sub> channel by binding to the N- and C-termini of GIRK1 (G protein-regulated inwardly rectifying K<sup>+<sup>) and GIRK4 subunits directly (Logothetis et al., 1987; Huang et al., 1995). The muscarinic acetylcholine K<sup>+</sup> channel (K<sub>Ach</sub>) is composed of GIRK1 and GIRK4 (Kir3.4, KCNJ5 gene) subunits in the atria and contributes to heart rate regulation in mammals (Wickman et al., 1998; Mark and Herlitze, 2000). During parasympathetic stimulation, K<sub>ach</sub> channels are activated, slowing the heart rate and the contractile force of the heart (Wickman et al., 1997). GIRK4 knockout mice lacked not only I<sub>Kach</sub> but also GIRK1 expression, reinforcing that GIRK4 plays a leading role in the expression and localization of GIRK1 to the cell membrane (Wickman et al., 1998; Kennedy et al., 1999). The K<sub>ach</sub> channel is rapidly and reversibly inhibited upon membrane stretch that allows mechano-electrical regulation of the atria (Li et al., 1998). Although GIRK4 is necessary for proper GIRK1 membrane expression, GIRK4 homotetramers have also been identified in the atria (Corey and Clapham, 1998; Bender et al., 2001). Regulation of the GIRK1/GIRK4 heterotetrameric channel occurs via phosphorylation by cyclic AMP-dependent protein kinase A (PKA) (upregulation) and via dephosphorylation by protein phosphatase 2A (PP2A) (downregulation) (Müllner et al., 2003).

Remarkably, the guanine nucleotide-binding protein subunits 2 and 5 (GNB2 and GNB5), which create the beta subunits of the G protein that interacts with GIRK1 and GIRK4, also play a role in SND. GNB5 variants have been reported in a patient with sinus bradycardia and cognitive disability (Lodder et al., 2016). A GNB2 variant was more strictly associated with cardiac conduction abnormalities resulting in SND and AV block (Stallmeyer et al., 2017).

**Human Studies**

Dobrev et al. (2001) compared 24 patients with chronic AF and aberrant atrial pacemaking function with 39 patients with normal sinus rhythm. The density of I<sub>Kach</sub> in patients with chronic AF was only 50% of the density of I<sub>Kach</sub> in the sinus rhythm group. GIRK4 mRNA expression was also decreased, potentially implicating reduced GIRK4 expression in increased atrial excitability (Dobrev et al., 2001). C171T and G810T variants in GIRK4 have now been identified as risk factors for lone paroxysmal AF in Chinese populations (Zhang et al., 2009). Further, A G387R heterozygous dominant variant in GIRK4 was found in a large Chinese kindred with clinical long QT syndrome (LQTS), a hereditary disorder that leads to SCD. In 2019, a GIRK4 variant (W1010C) was first identified in a three-generation family with SND. The W1010C variant in GIRK4 was interestingly resulted in increased I<sub>Kach</sub> expression (Dobrev et al., 2001). This gain-of-function variant caused an enhanced parasympathetic tone, causing familial SND and hyperpolarization of the pacemaker cells (Küf et al., 2019). Overall, loss-of-function variants seem to induce atrial arrhythmias, while gain-of-function variants may be implicated in SND.

**Mouse Models**

SAN pacemaker cells from Kcnj5<sup>−/−</sup> mice completely lacked I<sub>Kach</sub> and showed about 50% reduction in cholinergic regulation of heart rate (Wickman et al., 1998; Mesaric et al., 2013), further supporting the importance of the GIRK4 in parasympathetic modulation. In addition, Kcnj5 knockout mice had a 10% increase in resting heart rate with this decreased parasympathetic
tissue. The ability of these mice to recover to resting heart rate after stress (exercise or pharmacological stimulation) was significantly prolonged (Mesirca et al., 2013). Overall, complete lack of Kcnj5 expression attenuates the parasympathetic tone and results in an increased HR. Population-wide variants should be replicated in mice to further study the specifics of the mutations, and their effect on the functionality of the GIRK4 protein overall.

**Voltage-Gated Sodium Channel Alpha Subunit 5 (SCN5A)**

**Background**

Atrial, ventricular, and Purkinje myocyte depolarization, which causes a complete contraction of the heart, is initially regulated by the cardiac sodium channel. Na$_v$1.5 (the pore-forming, ion-conducting α-subunit of the cardiac sodium channel) is encoded by SCN5A, and variants in this gene have been implicated in a wide range of cardiac diseases such as Brugada syndrome, LQTS, AF, SND, dilated cardiomyopathy, and others (Wilde and Amin, 2018). Butters et al. (2010) analyzed electrophysiological mathematical models of SAN cells, 2D models of the intact SAN-atrium tissue, and actual recordings of activation patterns from isolated intact rabbit SAN-atrium tissue to elucidate the mechanism of Scn5a$^{+/−}$ variants on SAN function. Isolated SAN cells from adult rabbits harboring a heterozygous Scn5a variant display slower pacemaking rates in the peripheral cells, but not in the SAN central cells, yet 2D models show intact atrium-SAN tissue to have a decreased pacemaking rate overall with AP conduction issues, which may potentially cause SAN exit block and sinus arrest, as seen in SND (Butters et al., 2010).

**Human Studies**

A SCN5A variant associated with LQTS type 3 (LQT3) and Brugada syndrome was first identified in a large family in 2001. Individual carriers within the family that harbored the variant displayed a lower heart rate and experienced marked QT prolongation during episodes of bradyarrhythmia, which resulted in sudden death within the family (van den Berg et al., 2001). Further, SCN5A human variants (usually autosomal recessive inheritance patterns) were linked to SND (Benson et al., 2003), dilated cardiomyopathy, conduction disorders (McNair et al., 2004; Freyermuth et al., 2016; Yang et al., 2017), and infant death syndrome (Denti et al., 2018).

A loss-of-function E161K Na$_v$1.5 variant was identified in two unrelated individuals with family history of bradycardia, SND, conduction disease, and Brugada syndrome. The reduced functional Na$_v$1.5 protein expression caused atrial, ventricular, and SAN conduction slowing. Diastolic depolarization rate and upstroke velocity were both reduced in E161K computational models (Smits et al., 2005). An L1821fs/10 SCN5A variant causing a C-terminus truncation was identified in a 12-year-old male diagnosed with SND; and when expressed in HEK-293 cells, Na$_v$1.5 current density was decreased by 90% (Tan et al., 2007). The SND phenotype identified in SCN5A variants seems to generally come secondary to Brugada syndrome and LQTS3. For example, a loss-of-function E1784K variant was identified in 41 individuals, and 39% of individuals had SND, while nearly all of them (93%) had LQT3 (Makita et al., 2008). However, some SCN5A variants have shown an SND phenotype without the Brugada-type ST elevation (Nakajima et al., 2013; Wilders, 2018; Alkorashy et al., 2020). An I230T homozygous Na$_v$1.5 variant was found in four children with SND, yet heterozygous carriers of the variant showed normal conduction (Neu et al., 2010). Additionally, a case report found an R121W Na$_v$1.5 novel variant in an individual diagnosed with SND (Holst et al., 2010). Finally, D349N and D1790N autosomal recessive variants in Na$_v$1.5 were associated with pediatric sinus arrest and SND (Kodama et al., 2013).

**Mouse Models**

Mice heterozygous for functional Na$_v$1.5 (Scn5a$^{+/−}$) demonstrated bradycardia and SA block due to slowed pacemaker rates and slower SA conduction, particularly in larger peripheral SAN cells (Lei et al., 2005). Interestingly, Scn5a$^{+/−}$ mice showed sex-dependent effects on SAN functioning in older mice, particularly shown as PR interval prolongation (in old males), RR interval prolongation (longer in old males), QTc prolongation (similar across both genders), and T-wave prolongation (longer in old males) (Jeevaratnam et al., 2010). Scn5a$^{+/−}$ mice also had decreased heart rate variability, reduced SAN automaticity, slowed SA conduction, increased fibrosis, and increased fibroblasts as consequences of the decreased Na$_v$1.5 expression, particularly in older-age mice (Royer et al., 2005; Hao et al., 2011).

**Hyperpolarization Activated Cyclic Nucleotide Gated Potassium Channel 4 (HCN4)**

**Background**

The ability of the SAN cells to spontaneously initiate electrical impulse comes from the funny current ($I_f$) activation, a Na$^+/K^+$ depolarization current. Funny current channels have inward current at diastolic voltages that are then activated by membrane hyperpolarization (“membrane clock” hypothesis for pacemaker automaticity) via binding to the intracellular cAMP that can be modified by sympathetic and parasympathetic transmitters, therefore modulating the heart rate (DiFrancesco, 1993; Accili et al., 2002; DiFrancesco, 2010). $I_f$ may also be regulated by cAMP-activated PKA in the SAN (Liao et al., 2010). While there are four members of the HCN channel family, many studies proposed that only HCN2 and HCN4-based channels are expressed in the murine SAN, with HCN4-based channels having a higher level of expression, confirming the role of HCN4-based channels in driving cardiac pacemaker activity (Moosmang et al., 2001; Xiao et al., 2010). However, Fenske et al. (2013) found HCN1-based channels highly expressed in the SAN and reported HCN1-based channels as critical for the stabilization of the lead pacemaker region in mice. HCN4 channels and beta-2 adrenergic receptors (β2-AR) form a complex that is essential for HCN4 channel regulation (Greene et al., 2012). Interestingly, the expression of HCN2 and HCN4 channels was shown to decrease at the SAN and to increase in the atria and pulmonary vein in
older age in dogs, which could account for the disproportionately older population afflicted with SND (Li et al., 2014; Du et al., 2017).

Human Studies
In 2003, an HCN4 variant (573X) causing truncation of HCN4 C-terminus was first identified in a patient with SND, presented as sinus bradycardia and chronotropic incompetence (Schulze-Bahr et al., 2003). Importantly, familial sinus bradycardia linked to C-terminus truncation and loss of cAMP-dependent regulation of HCN4 was documented in 2010 (Schweizer et al., 2010). A D553N HCN4 missense variant was identified in an individual with recurring syncope, QT prolongation, polymorphic VT, and torsade de pointes. The individual variant, when transfected in COS7 cells, displayed a reduction in HCN4 expression at the cellular membrane, correlating the variant and the loss-of-function HCN4 with SND (Ueda et al., 2010). Sixteen family members carrying a G480R missense variant in HCN4 (autosomal dominant) experienced sinus bradycardia. Molecular studies showed reduced synthesis and trafficking of G480R variant HCN4 to the membrane (Nof et al., 2007). A G482R HCN4 variant was reported in a family presenting with bradycardia and left ventricular non-compaction cardiomyopathy (NCCM). Interestingly, this was one of the first studies to link HCN4-associated SND with structural abnormalities of the myocardium (Milano et al., 2014; Schweizer et al., 2014). Further, HCN4-R393H loss-of-function, c.1737+1G>T splice site, I1479V loss-of-function, A485E loss-of-function, R375C loss-of-function, and V759I loss-of-function variants have all been identified in individuals with SND and other related cardiac conduction disorders (Hategan et al., 2017; Ishikawa et al., 2017; Servatius et al., 2018; Alonso-Fernández-Gatta et al., 2020; Erlenhardt et al., 2020).

Mouse Models
Hcn2-deficient mice presented spontaneous absence seizures and abnormal cardiac sinus rhythm, implicating the HCN family in SAN function (Ludwig et al., 2003). Mice expressing the dominant negative 573X HCN4 isoform lacking cAMP regulation present with SAN bradycardia, similar to that observed in the SND individual (Alig et al., 2009). Conditional knockout of Hcn4 has produced variable outcome phenotype, from sinus pauses (Herrmann et al., 2007) to severe heart block and death (Baruscotti et al., 2011). A murine tamoxifen-inducible, cardiac-specific knockout model of exon 2 of Hcn4 channels showed remarkable bradycardia (50% reduction in heart rate), AV block, and death on day 5 on average. Importantly, I_F was reduced by about 70% in these mice (Baruscotti et al., 2011). A similar pattern of strong bradycardia and AV block was observed in mice expressing a dominant negative HCN4 subunit lacking channel conductance and completely lacking I_F. However, no mortality was observed in these mice, despite the presence of recurrent ventricular arrhythmia (Mesirca et al., 2014). Although HCN4 expression is essential for proper SAN functioning, extreme activation of I_F in cardiac cells can lead to initiation of ectopic foci, resulting in atrial and ventricular arrhythmias (Stieber et al., 2004). Morris et al. isolated atrial pacemaker cells from rats and found that overexpression of Hcn2 via adenovirus-mediated gene transfer resulted in pacing acceleration. This potentially implicates gain-of-function variants in HCN2 and HCN4 with increased pacemaker activity resulting in ectopic foci and cardiac arrhythmia (Morris et al., 2013). Interestingly, an Hcn1-deficient mouse model also showed bradycardia, slowed SAN conduction, sinus arrhythmia, and sinus pauses. This study suggests a role of HCN1 in SAN function and human SND (Fenske et al., 2013). Although the phenotypes for HCN4 variants in humans are varied, they nicely represent the various phenotypes in Hcn4-deficient mouse models.

Sodium/Calcium Exchanger 1 Precursor (SLC8A1)
Background
Spontaneous action potentials in SAN cells provide the primary pacemaking ability for the entire heart and are important for proper cardiac functioning. The cardiac Na+-Ca2+ exchanger (NCX1) plays an integral role in diastolic depolarizations that trigger these recurrent action potentials. Following diastolic SR Ca2+ release from ryanodine receptors, increased cytosolic Ca2+ causes an inward current via NCX, which accelerates late diastolic depolarization to the action potential threshold (“calcium clock” model for pacemaker automaticity) (Bogdanov et al., 2001; Lakatta et al., 2010). Further, NCX1 inactivation has the ability to completely halt SAN firing by generating intermittent burst firing induced by intracellular Ca2+ overload (Groenke et al., 2013; Torrente et al., 2015). NCX1 has 10 transmembrane helices and four ion-binding sites, one for Ca2+ and three for Na+ (Liao et al., 2012; Secondo et al., 2015). This NCX1 structural study agrees with current stoichiometric studies showing an exchange rate of mostly three Na+ to one Ca2+ (Bers and Ginsburg, 2007). NCX1 can function to facilitate either inward or outward current depending on the membrane potential (Bers, 2002). NCX1 is the predominant pathway for calcium extrusion in cardiomyocytes during resting membrane potential; the high extracellular sodium concentration allows NCX1 to exchange calcium out of the cell. Overall, NCX promotes myocytes to relax, therefore implicating a role of NCX1 in contractility. Further, spontaneous pacemaker release of Ca2+ by RyR2 activates the NCX1 on the SR membrane, which then pushes the cell to the minimum threshold for triggering an action potential (Shattuck et al., 2015).

Human Studies
Genetic variants in SLC8A1, the gene encoding NCX1, are associated with numerous electrocardiographic traits due to changes in calcium cycling. Polymorphisms in SLC8A1 across human populations were first associated with hypertension. Seven NCX1 polymorphisms with a high minor allele frequency of more than 4% were identified in 1,865 individuals, with 787 being hypertensive (Kokubo et al., 2004). Hong et al. (2014) found a single-nucleotide polymorphism on the SLC8A1 locus correlated with PR interval prolongation by a genome-wide association study using the Korea Association Resource database. Kim et al. (2012) also found a common SLC8A1 variant
associated with prolongation in QT interval, suggesting the predisposition of these populations to ventricular arrhythmias and SCD. Further, patients homozygous for allele rs13017968 in SLCA81 had higher rates of coronary artery abnormalities, predisposing these populations to Kawasaki disease. At present, there are no human data correlating mutations on SLCA81 with SND. However, animal models showed major phenotypes of SND when experiencing loss of NCX1 (Gao et al., 2013; Groenke et al., 2013; Torrente et al., 2017).

**Mouse Models**

The role of NCX1 in calcium cycling and its importance to the SAN have been studied through multiple Slc8a1-null mouse models. A global Slc8a1 knockout resulted in abnormal myofibrillar organization and severe electrical defects that caused embryonic lethality (Fu et al., 2010); therefore, cardiac-specific knockouts are required to study the role of NCX1 specifically in the heart. Gao et al. (2013) used a global myocardial and SAN-targeted knockout of Slc8a1 to study the role of NCX1 in pacemaker activity. Although isolated SAN cells showed similar basal contractility rates in SLCA81 knockout versus WT mice, Slc8a1 knockout mice showed the inability to respond to isoproterenol, implicating a role of NCX1 in the sympathetic response of the heart (Gao et al., 2013). Several studies have established the role of NCX1 in calcium efflux; however, the role of NCX1 in triggering an action potential is still not completely understood. Groenke et al. (2013) showed pacemaker activity to be completely ablated in an atrial-specific Slc8a1 knockout mouse model. The Slc8a1 atrial specific knockout mouse also lacked P waves and had arrhythmic depolarizations in the SAN. Although SAN automaticity still occurred without NCX1, the automaticity came in bursts similar to tachycardia–bradycardia syndrome and SND (Torrente et al., 2015). Torrente et al. (2015) further used their atrial-specific Slc8a1 knockout and found severe cellular Ca²⁺ accumulation during SA nodal pacemaker activity, leading to intermittent hyperactivation of small conductance K⁺ (SK) channels, subsequently resulting in arrhythmias. These data identified the potential influence of intracellular Ca²⁺ on SK channels and overall SAN repolarization and signified SK channels as potential therapeutic targets for SAN dysfunction if presented alongside Ca²⁺ cycling issues (Torrente et al., 2017).

**Ankyrin-2 (ANK2)**

**Background**

Ankyrin-B (AnkB, encoded by ANK2) is a membrane adapter protein critical in the recruitment, organization, and stabilization of the ion channels and transporters underlying the excitation–contraction coupling, particularly in the SAN. Loss-of-function variants in ANK2 are associated with a complex cardiac phenotype including heart rate variability, CPVT, conduction defects, AF, sinus node bradycardia, SCD, and, recently, arrhythmogenic cardiomyopathy (Roberts et al., 2019).

**Human Studies**

Two families characterized with severe SND were found to have ANK2 allele variants, making AnkB the first non-ion channel protein associated with human SND (Curran and Mohler, 2011). Interestingly, individuals with AF have reduced levels of AnkB expression and increased levels of miR-34a (a microRNA associated with cardiac fibrosis). Of note, the 3′ untranslated region of ANK2 also contains the binding site to miR-34a, implicating a potential role of miR-34a in the electrical remodeling of the atria and in the regulation of AnkB expression (Zhu et al., 2018). Although the culmination of cardiac AnkB studies implicate loss-of-function ANK2 variants in numerous cardiac diseases, the lack of family history in many of these cases, and overall incomplete penetrance of AnkB-associated disease, strongly implies that additional genetic and/or environmental factors must be involved in the development of the severe “AnkB syndrome” phenotype. Notably, intense endurance exercise or other genetic variants likely play a role in the development of cardiac disease associated with loss-of-function ANK2 variants (Roberts et al., 2019).

**Myosin Heavy Chain 6 (MYH6)**

MYH6 encodes the alpha myosin heavy chain subunit of myosin (MHC-α), a major component of the sarcomere—a necessary component of muscle fiber for proper contraction in the heart (Epp et al., 1993; Squire, 1997). An MHC-α R721W missense variant has been identified in Icelandic populations (0.38% allelic frequency) and is associated with SND, with 50% of the carriers for this variant being diagnosed with SND. Carriers of the variant that were not diagnosed with SND still showed reduced heart rate and PR interval prolongation (Holm et al., 2011). Interestingly, another R654W heterozygous MYH-α variant was identified in an Australian family with severe yet diverse cardiac arrhythmias, including SND and cardiac arrest due to ventricular fibrillation, resulting in SCD or SND (Lam et al., 2015).

**Lamin A (LMNA)**

Nuclear lamins (lamins A, B1, and B2) are the major components of nuclear lamina, which plays a vital structural role in the nuclear envelope (Mounkes et al., 2001). LMNA variants are associated with numerous cardiac conditions, particularly dilated...
Although no population-wide variant in \textit{LMNA} by nonsense-mediated mRNA decay (Zaragoza et al., 2016). As aberrant mRNA from the mutant allele would likely decay by nonsense-mediated mRNA decay (Zaragoza et al., 2016). Although no population-wide variant in \textit{LMNA} by nonsense-mediated mRNA decay (Zaragoza et al., 2016). As aberrant mRNA from the mutant allele would likely decay by nonsense-mediated mRNA decay (Zaragoza et al., 2016). Although no population-wide variant in \textit{LMNA} by nonsense-mediated mRNA decay (Zaragoza et al., 2016). As aberrant mRNA from the mutant allele would likely decay by nonsense-mediated mRNA decay (Zaragoza et al., 2016). Although no population-wide variant in \textit{LMNA} by nonsense-mediated mRNA decay (Zaragoza et al., 2016). As aberrant mRNA from the mutant allele would likely decay by nonsense-mediated mRNA decay (Zaragoza et al., 2016).

### L-Type Calcium Channel Subunit Ca\textsubscript{v}1.3 (CACNA1D)

#### Background

Ca\textsubscript{v}1.2 (alpha1C) and Ca\textsubscript{v}1.3 (alpha1D) subunits make up the cardiac L-type voltage-activated calcium channel (Matthes et al., 2004). Ca\textsubscript{v}1.3 is expressed mostly in the SAN, AV node, and atrial myocytes (Mangoni et al., 2003). In the SAN, Ca\textsubscript{v}1.3 plays a major role in pacemaker activity by driving inward current during diastolic depolarization and regulating diastolic SR Ca\textsuperscript{2+} release (Mangoni et al., 2003; Torrente et al., 2016).

T-type calcium channels are composed of three subunits, Ca\textsubscript{v}3.1, Ca\textsubscript{v}3.2, and Ca\textsubscript{v}3.3, which are encoded by three genes: CACNA1G, CACNA1H, and CACNA1I. The three Ca\textsubscript{v}3 subunits are responsible for the generation of T-type/low-voltage activated calcium currents (T-current) (Huc et al., 2009). Variants in CACNA1H and CACNA1G were identified in epileptic patients (Chen et al., 2003; Singh et al., 2007). Interestingly, earlier studies reported the expression of T-type calcium current (\textit{I}_{\text{Ca,T}}) in pacemaker cells and proposed its contribution to pacemaking function in SAN cells (Hagiwara et al., 1988). Later, Mangoni et al. (2006) demonstrated the direct contribution of Ca\textsubscript{v}3.1 channels to pacemaking and cardiac conduction through genetic ablation of Ca\textsubscript{v}3.1 channels in mice. The genetic inactivation of Ca\textsubscript{v}3.1 channels caused a slowing of pacemaking function through a reduction of the slope of the diastolic depolarization in SAN cells (Mangoni et al., 2006).

#### Mouse Models

\textit{cacna1d}\textsuperscript{-/-} mice have severe SND and AV second-degree block (Platzer et al., 2000; Zhang et al., 2002; Mesirca et al., 2016). In addition, Ca\textsubscript{v}1.3 also plays an important role in calcium homeostasis in the ear, as \textit{cacna1d}\textsuperscript{-/-} mice experience deafness along with SND (Chu et al., 2007). The phenotype of SND-associated deafness was later identified in two related families with a Ca\textsubscript{v}1.3 variant (G403_V404insG). This rare combination of symptoms is termed sinoatrial node dysfunction and deafness (SANDD) (Baig et al., 2011). This CACNA1D variant was also identified in four additional families with SANDD, along with an A376V missense variant (Liaqat et al., 2019).

#### Short-Statue Homeobox 2 (SHOX2)

#### Background

Short-stature homeobox 2 is a transcription factor encoded by \textit{SHOX2}. \textit{SHOX2} is essential for the proper development of the SAN (Puskaric et al., 2010). \textit{Shox2}\textsuperscript{-/-} mice are embryonically lethal due to lack of SAN development, but \textit{Shox2}\textsuperscript{+/+} zebrafish survive with bradycardia (Hoffmann et al., 2013). Mutations in genes encoding subunits and/or proteins directly involved or being components of the cardiac conduction system were linked to rhythm abnormalities (Frank et al., 2012). Interestingly, mutations in TBX genes such as \textit{TBX3} and \textit{TBX5} were associated with conduction dysfunction (Wolf and Berul, 2006; Christoffels et al., 2010). Notably, the transcription factor \textit{NKX2-5} has also been identified as a regulator of heart rate, and several variants linked to \textit{NKX2-5} were shown to cause heart rate variability and increased incidence of AF (den Hoed et al., 2013; Huang et al., 2013). Finally, transcriptome analysis of the SAN cells demonstrated a conserved genetic program between mouse and human cells including \textit{TBX3}, \textit{SHOX2}, \textit{ISL1}, \textit{BMP}, and \textit{NOTCH} signaling components (van Eif et al., 2019).

#### Human Studies

Loss-of-function variants in \textit{SHOX2} cause early-onset and familial AF (Hoffmann et al., 2016; Li et al., 2018). A heterozygous missense P33R \textit{SHOX2} variant was also identified recently in a patient with SND (Hoffmann et al., 2019). Screening should continue for \textit{SHOX2} variants in patients with SND, as this gene has only been recently implicated in SAN function in humans.

#### Transient Receptor Potential Cation Channel Subfamily C Member 3 (TRPC3)

#### Background

TRPC channels (non-selective Ca\textsuperscript{2+}-permeable cation channels) are thought to play an important role in store-operated Ca\textsuperscript{2+} entry (SOCE), described as Ca\textsuperscript{2+} influx to the sarcolemma after Ca\textsuperscript{2+} depletion, although the pathway is currently poorly understood (Wu et al., 2004; Ju et al., 2007). TRPC channels are activated when diacylglycerol is released from the plasma membrane via agonist binding to G protein-coupled receptors (Wu et al., 2004). All TRPC subtypes (1–7) are expressed in the SAN except subtype 5, but TRPC3 is the only subtype expressed on the membrane surface of the pacemaker cells (Ju et al., 2007).

#### Human Studies

Human studies about TRPC channels are greatly needed. Current TRPC3 research is almost completely focused on animal models. One study identified marked increases of leukocyte TRP channel mRNA in 47 patients with non-valvular AF, implicating upregulation of TRPC channels in AF (Düzen et al., 2017).

#### Mouse Models

\textit{Trpc3}\textsuperscript{-/-} mice treated with angiotensin II showed increased spontaneous action potentials (Qi et al., 2015). \textit{Trpc3} specifically has been shown to increase local Ca\textsuperscript{2+} release (LCR) and NCX current (\textit{I}_{\text{NCX}}) in mouse embryonic stem cell-derived cardiomyocytes, which resulted in increased spontaneous action potentials (Qi et al., 2016). Both of these studies support the hypothesis that Ca\textsuperscript{2+} entry via TRPC3 is a pro-arrhythmic pathway, which potentially causes increased pacemaker activity. TRPC3 is
upregulated in AF patients and AF animal models (Harada et al., 2012). Finally, TRPC-3 channel upregulation also has been shown to cause an increased accumulation of collagen consistent with atrial fibrosis in mice (Han et al., 2020). FK506-binding protein 52 (FKBP52 or KBP4) has been identified as an interaction partner of TRPC3 and may be an important player in TRPC3-related therapy going forward. Downregulation of FKBP52 induced a Trpc3-dependent hypertrophic response in neonatal rat cardiomyocytes (Bandleon et al., 2019). Upregulation of FKBP52 may be able to reduce activity of the pro-arrhythmic TRPC3 pathway, but more studies are needed to understand this interaction. TRPC also has potential as a target for cardiac fibrogenesis (Yue et al., 2013). TRPC3 has also been implicated in cardiac hypertrophy, and gene deletion of Trpc3 and Trpc6 in mice protected against pressure-induced cardiac remodeling (Seo et al., 2014).

**Stromal Interaction Molecule 1 (STIM1)**

Background

SOCE is an important pathway for Ca^{2+} reentry after calcium depletion, particularly in cardiac pacemaker cells. STIM1 is an endoplasmic reticulum Ca^{2+} sensor essential to the SOCE pathway, implicating STIM1 in the calcium clock of SAN cells. Following Ca^{2+} depletion in the SAN cells, STIM1 localizes to the cell periphery, along with calcium release-activated calcium channel protein 1 (Orai1) (Liu et al., 2015). STIM1 and Orai1 channels are selectively expressed in SAN cells, and STIM1 is essential in SAN functioning.

Mouse Models

Stim1 cardiac knockout mice showed SR calcium store depletion in SAN cells, resulting in SAN dysfunction (Zhang et al., 2015). STIM1-deficient mice experienced slowed heart rate after stimulation, sinus arrest, and extreme autonomic response to cholinergic signaling. Further, they also showed reduction in L-type Ca^{2+} current and enhanced NCX activity, linking STIM1 to more regulatory pathways in Ca^{2+} cycling in SAN cells than previously anticipated (Zhang et al., 2015). Beyond the SAN, STIM1 has been found to play an essential role in interatrial conduction via its expression in sinus cardiomyocytes from the SAN to the coronary sinus. Deletion of Stim1 from coronary sinus cardiomyocytes slowed conduction across the atria and increased susceptibility to atrial arrhythmias in Stim1 cardiac-specific knockout mice (Zhang et al., 2020).

**Potassium Two Pore Domain Channel Subfamily K Member 2 (KCNK2)**

Background

TREK-1, or K2p2.1 (encoded by KCNK2 gene), is a K^{+} channel with four transmembrane segments and two pore domains that are activated via membrane stretch and arachidonic acid, among other mechanisms (Fink et al., 1996; Lesage and Lazdunski, 1998; Maingret et al., 1999). K^{+} channels in cardiac tissue open to cause hyperpolarization and close during depolarization, therefore playing a crucial role in selecting the duration of an action potential (Snyders, 1999). Beyond membrane stretch and arachidonic acid, TREK-1 is regulated by polyunsaturated fatty acids, temperature, receptor-coupled second messenger systems, volatile anesthetics, neuroprotectant agents, and selective serotonin reuptake inhibitors (Goonetilleke and Quayle, 2012).

**Transient Receptor Potential Melastatin 4 (TRPM4)**

TRPM4 is a selective monovalent cation channel that allows flow of Na^{+}, K^{+}, Cs^{+}, and Li^{+} (Guinamard et al., 2010), which is activated by intracellular Ca^{2+} (Launay et al., 2002). A study in 160 unrelated probands identified multiple TRPM4 variants associated with right-bundle branch block and isolated AV block, signifying the role of TRPM4 in cardiac conduction. Surprisingly, none of the patients with SND harbored TRPM4 variants (Stallmeyer et al., 2012). However, Trpm4 is expressed in mouse SAN cells and is potentially implicated in heart rate rhythm regulation (Demion et al., 2007). Further, TRPM4 inhibition showed the ability to reduce the action potential rate by modulation of Ca^{2+}-activated non-selective cation current in isolated mammalian right atrial cells (Hof et al., 2013). Therefore, patients with SND should continue to be screened for TRPM4 variants in the future considering the involvement of TRPM4 involvement in cardiac conduction.

**CLINICAL MANAGEMENT OF SINOATRIAL NODE DYSFUNCTION**

Pharmacologic Approach

Many genetic forms of SND are chronic, yet symptoms are mild. Acute SND, however, can appear secondary to another condition, procedure, or disease, as mentioned previously. Acute SND presented as bradycardia or bradycardia/tachycardia syndrome is dangerous and can be managed with a small selection of pharmacological agents. Therapeutic control of SND is ideal because of the reduced cost and avoidance of surgical intervention. Current pharmacological options for acute management of bradycardia associated with SND include atropine, isoproterenol, aminophylline, or theophylline according to the American College of Cardiology/American Heart Association/Heart Rhythm Society (ACC/AHA/HRS) updated guidelines in 2019 (Kusumoto et al., 2019).

Atropine may successfully reverse an acute SND condition to a normal sinus rhythm (Schweitzer and Mark, 1980). Atropine shortens the sinus cycle length and the sinus recovery time of the SAN, therefore potentially treating bradycardia...
Acute bradycardia following heart transplant is a common outcome. As mentioned earlier, atropine is a poor choice for bradycardia therapy after heart transplant due to the potential for complete AV block and sinus arrest (Bernoheim et al., 2004). In this scenario, theophylline has displayed the ability to reverse bradycardia in the majority of post-transplant patients, averting the need for pacemaker implantation (Bertolet et al., 1996; Kertesz et al., 2003). After heart transplant, relative bradycardia is often considered as a heart rate of less than 80 bpm, as postoperative patients have higher hemodynamic demand. The lack of parasympathetic input in a donor heart often results in SND (Woo et al., 2008). Bertolet et al. (1996) showed the mean heart rate of 29 patients suffering from bradycardia post-heart transplant was improved from 62 ± 7 to 89 ± 10 bpm when treated with theophylline (300 mg intravenous, 474 ± 99 mg/day orally) after 24 h of treatment (Bertolet et al., 1996). Aminophylline has also seen some supportive data, but theophylline is more supported across the literature (Kusumoto et al., 2019). Theophylline shows some promise in treating chronic SND as well (Alboni et al., 1991; Saito et al., 1993), but theophylline has numerous situational restrictions that prevent it from becoming the gold standard for chronic SND therapy. Theophylline should specifically be avoided in cases of bradycardia/tachycardia syndrome or in patients experiencing any ventricular ectopy (Ling and Crouch, 1998).

Beyond therapeutic purposes, atropine and isoproterenol can be used to evaluate the severity of asymptomatic sinus bradycardia. Interestingly, the chronotropic response to atropine or isoproterenol can identify patients in need of preventative pacemaker implantations (Vavetsi et al., 2008).

Finally, a study of 192 patients with SND showed that cilostazol was able to prevent permanent pacemaker implantation. Only 20.4% of patients receiving cilostazol needed a pacemaker implant, while 55.8% of patients not receiving cilostazol required a pacemaker. By increasing heart rate, cilostazol shows promise to be the first long-term pharmacological therapeutic for SND as many of the pharmacological options discussed earlier are only effective short-term and have negative long-term side effects (Sonoura et al., 2019). With Sonoura et al. (2019) publishing this study in 2019, long-term studies showing longer-term survivability over 5 or 10 years in patients using cilostazol would further support the use of this drug to manage SND.

**Temporary Pacemaking**

When pharmacological therapies fail, temporary pacemakers are an alternative approach and can be implemented via a multitude of methods for care of acute bradycardia. Transcutaneous cardiac pacing is an external pacing method (prevents the risk of infection and other surgical complications) and is commonly used in response to patients with life-threatening bradycardia (Dalsey et al., 1984). Transcutaneous pacing does, however, show success in reverting bradycardia in patients not in cardiac arrest too (Clinton et al., 1985). Transvenous pacemaker therapy is another temporary pacemaker option. Transvenous pacemakers can be implanted in the internal and external jugular, subclavian, brachial, and femoral veins, although the right internal jugular vein is the most preferred location for best access to the right ventricle (Gammage, 2000). The early studies of transvenous pacemaker therapy showed very high rates of pacemaker malfunction (as high as 43%) (Lumia and Rios, 1973). Even today, approaches for temporary transvenous cardiac pacing vary widely, so outcomes are difficult to compare across studies (Diemberger et al., 2020). However, a study in 2018 across more than 360,000 patients found complications for modern-day temporary transvenous pacing to be only about 4%, but with 37.9% of patients still needing permanent pacemaker implantation (Metkus et al., 2019). Overall, the ACC/AHA/HRS guidelines recommend transcutaneous pacing over transvenous pacemaker therapy due to the complications of the transvenous method when a patient is hemodynamically unstable and critically ill due to bradycardia (Kusumoto et al., 2019). This suggestion may be changed in the future with large group studies showing the success of transvenous pacing.

**Permanent Pacemaker Implant**

In general, a permanent pacemaker implant should be avoided if possible to avert the potential procedural complications related to the implant and long-term management of the implant. Unfortunately, permanent pacemaker implantation may be necessary to prevent severe and life-threatening arrhythmias and should be considered with severe symptomatic sinus bradycardia and bradycardia/tachycardia syndrome (Kusumoto et al., 2019). For example, permanent pacemaking after Fontan operation, a surgery for children with only a single functional ventricle, often results in SND and requires permanent atrial pacing in 13–16% of patients (Cohen et al., 1998; Takahashi et al., 2009). Further, patients presenting with AF and significant atrial fibrosis may
require a permanent pacemaker implant (Akoum et al., 2012). Atrial flutter, associated with SND, can also be used to predict the need for pacemaker implantation. In 211 patients following ablation, an atrial flutter length of more than 273 ms predicted the need for a pacemaker implantation (Sairaku et al., 2012).

**Atrial Ganglionated Plexus Modification**

Permanent pacemaker implantation is still the current therapeutic suggestion for long-term sinus bradycardia by the AHA (Tracy et al., 2013). However, younger SND patients will on average require a longer life from the permanent pacemaker and, therefore, require lead and generator replacements, which increase risk of infection and surgical complications (Qin et al., 2017). Notably, the sympathetic and parasympathetic nervous systems modulate sinus rhythm via the atrial ganglionated plexi (GP). Zhao et al. (2015) used endocardial radiofrequency ablation in the GP to prevent parasympathetic nervous system hyperactivity and successfully increased resting heart rate in young SND patients.

**FUTURE IMPLICATIONS**

SND is a disorder caused by the inability of the heart to perform its pacemaking function (Semelka et al., 2013). It is manifested clinically as sinus bradycardia, sinus pause, or alternating bradycardia and tachyarrhythmias. Importantly, patients report chronotropic incompetence in response to stress or exercise (Semelka et al., 2013). Permanent pacemaker implantation is only recommended for symptomatic patients. However, there is no existing therapy to reverse the primary genetic cause in patients diagnosed with primary or chronic SND. Calcium-activated potassium channels (SK4) have been implicated in the automaticity of cardiomyocytes, yet Haron-Khun et al. (2017) showed that TRAM-34 (a selective blocker of SK4 channels) successfully reduced DADs and calcium transients in human-induced pluripotent stem cell-derived cardiomyocytes (hiPSC-CMs) from patients with CPVT2 from a CASQ2-D307H variant. When mice with the CASQ2-D307H knock in variant were injected with TRAM-34, electrocardiographic recordings showed reduced arrhythmias (and reduced severity of arrhythmias) at rest and during exercise. This was also shown in CASQ2 KO mice. TRAM-34 and clotrimazole (SK4 inhibitors) therefore hold great therapeutic promise for human populations with loss-of-function CASQ2 variants (Haron-Khun et al., 2017).

Non-functional cardiac funny current (I_f) causes unusual Ca^{2+} handling as previously discussed, therefore disrupting the pacemaker activity by the sinus node. Remarkably, Mesirca et al. (2014) showed how cardiac funny current-deficient mice have impulse generation and conduction defects, which can be rescued with genetic deletion of cardiac muscarinic G-protein-activated channels (GIRK4). Although HCN4 and GIRK4 loss-of-function variants have each been implicated in SND, the combination of silencing both genes seems to repair severe cardiac arrhythmia phenotype of SND associated with AV block and ventricular arrhythmia (Mesirca et al., 2014). Interestingly, follow-up work has indicated that genetic deletion of GIRK4 also rescues conduction defects in model mice of Ca_{v}1.3-mediated SND (Mesirca et al., 2016). Furthermore, rescuing of SND in mice models carrying dysfunction in HCN4, Na_{v}1.5, and Ca_{v}1.3 channels can be mimicked by acute administration of the I_{KACH} blocker tertapiqin-Q (Bidaud et al., 2020). Importantly, the ability of pharmacologic inhibition of I_{KACH} to improve SAN function has been demonstrated recently in human SANs with history of SND (Li et al., 2017). Finally, silencing of GIRK4 expression in human atrial myocytes was shown to efficiently decrease I_{KACH} densities and therefore is a great potential tool for treating arrhythmia (Liu et al., 2009). In conclusion, data from murine models and human SAN tissues indicate that gene therapy or pharmacologic strategy targeting GIRK4 channels can constitute an important future direction for clinical management of SND.

**HCN4 lentiviral gene transfer** has interestingly shown the bioengineering potential to allow for pacemaker cell therapy. Transducing HCN4 revived autonomous pacemaking and increased responsiveness to autonomic regulation in HCN4-transduced myocytes (Boink et al., 2008). Further, myocyte enhancer factor-2 (MEF2) and activator protein-1 (AP1), with binding sequences located on conserved non-coding sequence 13 (CN13), are involved in HCN4 enhancement via the HCN4 promoter (Kuratomi et al., 2009) and could be used to upregulate HCN4 to promote pacemaking activity.

In summary, SND is a disorder that more commonly affects the elderly population and impacts the pacemaking function of the heart resulting in arrhythmia and chronotropic incompetence. The complexity of SND is partially attributed to the complexity of genetic abnormalities and partially attributed to gene pleiotropy. Many variants of SND-associated genes can exhibit multiple unrelated phenotypic traits. Currently, clinical management of SND patients is restricted to the treatment or relief of arrhythmia symptoms. However, there is no widely available therapeutic option that targets or reverses the primary genetic cause in patients with chronic SND. Understanding the complexity of genetics that contribute to disease progression is critical to developing new therapeutic strategies for this complex, life-threatening disorder.

**AUTHOR CONTRIBUTIONS**

All authors listed have made a substantial, direct and intellectual contribution to the work, and approved it for publication.

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REFERENCES

Abe, K., Machida, T., Sumitomo, N., Yamamoto, H., Ohkubo, K., Watanahe, I., et al. (2014). Sodium channelopathy underlying familial sick sinus syndrome with early onset and predominantly male characteristics. Circ. Arrhythm. Electrophysiol. 7, 511–517. doi: 10.1161/CIRCEP.113.001340

Accili, E. A., Proenza, C., Baruscotti, M., and DiFrancesco, D. (2002). From funny current to HCN channels: 20 years of excitement. News Physiol. Sci. 17, 32–37. doi: 10.1152/physiolsyn.2002.17.1.32

Akoum, N., McGann, C., Vergara, G., Badger, T., Ranjan, R., Mahnkopf, C., et al. (2012). Atrial fibrillation quantified using late gadolinium enhancement MRI is associated with sinus node dysfunction requiring pacemaker implant. J. Cardiovasc. Electrophysiol. 23, 44–50. doi: 10.1111/j.1540-8167.2011.02140.x

Alboni, P., Ratto, B., Cappato, R., Rossi, P., Gatto, E., and Antonioli, G. E. (1991). Clinical effects of oral theophylline in sick sinus syndrome. Am. Heart J. 122, 1361–1367. doi: 10.1016/0002-8703(91)90578-6

Ali, J., Marger, L., Mesirca, P., Ehmkhe, H., Mangoni, M. E., and Isbrandt, D. (2009). Control of heart rate by cAMP sensitivity of HCN channels. Proc. Natl. Acad. Sci. U.S.A. 106, 12189–12194. doi: 10.1073/pnas.081032106

Alings, A. M., Abbas, R. F., and Bouman, L. N. (1995). Age-related changes in current to HCN channels: 20 years of excitation. J. Physiol. 491 Pt 3, 617–633. doi: 10.1113/jphysiol.1995.sp021042

Allan, J. P., Edin, H., Svedmyr, I., and Karlsson, A. E. (2009). The human cardiac SR Ca(2+)-ATPase: a dual role in metazoan calcium homeostasis. Cell Calcium 46, 109–122. doi: 10.1016/j.ceca.2009.08.004

Alig, J., Marger, L., Mesirca, P., Ehmkhe, H., Mangoni, M. E., and Isbrandt, D. (2009). Control of heart rate by cAMP sensitivity of HCN channels. Proc. Natl. Acad. Sci. U.S.A. 106, 12189–12194. doi: 10.1073/pnas.081032106

Ali, J., Marger, L., Mesirca, P., Ehmkhe, H., Mangoni, M. E., and Isbrandt, D. (2009). Control of heart rate by cAMP sensitivity of HCN channels. Proc. Natl. Acad. Sci. U.S.A. 106, 12189–12194. doi: 10.1073/pnas.081032106
impaired cardiac pacemaking in sick sinus syndrome. *Circ. Res.* 107, 126–137. doi: 10.1161/circresaha.111.300076

Chen, Y., Lu, J., Pan, H., Zhang, Y., Wu, H., Xu, K., et al. (2003). Association of mutation of the gene encoding sarcolemmal Ca2+-ATPase with cardiac conduction and rhythm disorders. *J. Emerg. Med.* 2, 155–162. doi: 10.1016/j.jemermed.2002.06.010

Corey, S., and Clapham, D. E. (1998). Identification of native atrial G-protein-regulated inwardly rectifying K+ (GIRK4) channel homomultimers. *J. Biol. Chem.* 273, 27499–27504. doi: 10.1074/jbc.273.42.27499

Cossù, S. F., Rothman, S. A., Chmielewski, I. L., Hsia, H. H., Vogel, R. L., Miller, C. J., and Mohler, P. J. (2011). Coordinating electrical activity of the heart: a tricky case of pediatric overlap syndrome? *Front. Physiol.* 2, 52. doi: 10.3389/fphys.2015.00037

Curran, J., and Mohler, P. J. (2011). Coordinating electrical activity of the heart: ankyrin polypeptides in human cardiac disease. *Expert Opin. Ther. Targets* 15, 789–801. doi: 10.1517/14728222.2011.575363

Dadlani, R., Challam, K., Garg, A., and Hegde, A. S. (2010). Can bradycardia pose a threat in patients with sinus nodal dysfunction? *Am. J. Cardiol.* 105, 183–195. doi: 10.1016/j.amjcard.2009.12.010

Dadlani, R., Challam, K., Garg, A., and Hegde, A. S. (2010). Can bradycardia pose a threat in patients with sinus nodal dysfunction? *Am. J. Cardiol.* 105, 183–195. doi: 10.1016/j.amjcard.2009.12.010

Dentl, F., Bentzen, B. H., Wojciak, J., Thomsen, N. M., Scheiman, M., and Schmitt, N. (2018). Multiple genetic variations in sodium channel subunits in a case of sudden infant death syndrome. *Pacing Clin. Electrophysiol.* 41, 620–626. doi: 10.1111/pcc.13528

Dharmawan, T., Nakajima, T., Ohno, S., Izuka, T., Tamura, S., Kaneko, Y., et al. (2019). Identification of a novel exon3 deletion of RYR2 in a family with catecholaminergic polymorphic ventricular tachycardia. *Ann. Noninvasive Electrocardiol.* 24:e12623. doi: 10.1111/anec.12623

Dhingra, R. C., Amit, Y. L. F., Wyndham, C., Denes, P., Wu, D., Miller, R. H., et al. (1976). Electrophysiological effects of atropine on sinus node and atrium in patients with sinus nodal dysfunction. *Am. J. Cardiol.* 38, 848–855. doi: 10.1001/0002-9149(76)90797-9

Dieringer, I., Massaro, G., Rossillo, A., Chieffo, E., Dugo, D., Guarracini, F., et al. (2020). Temporary transvenous cardiac pacing: a survey on current practice. *J. Cardiovasc. Med.* 21, 420–427. doi: 10.2459/jcm.0000000000000958

D’Inverno (1993). Pacemaker mechanisms in cardiac tissue. *Annu. Rev. Physiol.* 55, 455–472. doi: 10.1146/annurev.physiol.55.030193.002323

DiFrancesco, D. (2010). The role of the funny current in pacemaker activity. *Circ. Res.* 106, 434–446. doi: 10.1161/circresaha.109.208041

Dobrev, D., Graf, E., Wettwer, E., Himmel, H. M., Håla, O., Doerfel, C., et al. (2007). Molecular basis of downregulation of G-protein-coupled inward rectifying K+ (current IK(Ach)) in chronic human atrial fibrillation: decrease in GIRK4 mRNA correlates with reduced I(ACh) and muscarinic receptor-mediated shortening of action potentials. *Circulation* 104, 2551–2557. doi: 10.1161/hc4601.099466

Dobrzynski, H., Boyett, M. R., and Anderson, R. H. (2007). New insights into pacemaker activity: promoting understanding of sick sinus syndrome. *Circulation* 115, 1921–1932. doi: 10.1161/circulationaha.106.616011

Domingo, D., Neco, P., Fernández-Pons, E., Zissimopoulos, S., Molina, P., Olagué, J., et al. (2015). Non-ventricular, clinical, and functional features of the RyR2(R420Q) mutation causing catecholaminergic polymorphic ventricular tachycardia. *Rev. Esp. Cardiol.* 68, 398–407. doi: 10.1016/j.recesp.2014.04.023

Du, J., Deng, S., Pu, D., Liu, Y., Xiao, J., and She, Q. (2017). Age-dependent down-regulation of hyperpolarization-activated cyclic nucleotide-gated channel 4 causes deterioration of canine sinoatrial node function. *Acta Biochim. Biophys. Sin.* 49, 400–408. doi: 10.1093/abbs/gmx026

Düzen, I. V., Yavuz, F., Vuruskun, E., Saracoglu, E., Poyraz, F., Göksuluk, H., et al. (2017). Leukocyte TRP channel gene expressions in patients with non-valvular atrial fibrillation. *Sci. Rep.* 7:7292. doi: 10.1038/s41598-017-10039-0

Epp, T. A., Dixon, I. M., Wang, H.-Y., Sole, M. J., and Liew, C.-C. (1993). Structural organization of the human cardiac α-myosin heavy chain gene (MYH6). *Genomics* 18, 505–509.

Erlenhardt, N., Kletke, O., Wohlforth, F., Komadowski, M. A., Clasen, L., Makimoto, H., et al. (2020). Disease-associated HCN4 V759I variant is not sufficient to impair cardiac pacemaking. *Pflugers Arch.* 472, 1733–1742. doi: 10.1007/s00424-020-02481-3

Faggioni, M., Hwang, H. S., van der Werf, C., Nederend, I., Kannankeril, P. J., Wilde, A. A., et al. (2013). Accelerated sinus rhythm prevents catecholaminergic polymorphic ventricular tachycardia in mice and in patients. *Circ. Res.* 112, 689–697. doi: 10.1161/circresaha.111.300076

Faggioni, M., Savio-Galimberti, E., Venkataraman, R., Hwang, H. S., Kannankeril, P. J., Darbar, D., et al. (2014). Suppression of spontaneous ca elevations prevents atrial fibrillation in casequestrin 2-null hearts. *Circ. Arrhythm. Electrophysiol.* 7, 313–320. doi: 10.1161/circep.113.000994

Fedele, L., and Brand, T. (2020). The intrinsic cardiac nervous system and its role in cardiac pacemaking and conduction. *J. Cardiovasc. Dev. Dis.* 7:54. doi: 10.3390/jcd7040054

Fenske, S., Krause, C., Hassan, S. I., Becirovic, E., Auer, F., Bernard, R., et al. (2013). Sick sinus syndrome in HCN1-deficient mice. *Circulation* 128, 2585–2594. doi: 10.1161/circulationaha.113.003712

Fink, M., Duprat, P., Lesage, F., Reyes, R., Romey, G., Heurtéaux, C., et al. (1996). Cloning, functional expression and brain localization of a novel unconventional outward rectifier K+ channel. *EMBO J.* 15, 6854–6862.

Frank, D. U., Carter, K. L., Thomas, K. R., Burr, R. M., Bakker, M. L., Coetzee, W. A., et al. (2012). Lethal arrhythmias in Tbx3-deficient mice reveal extreme dosage sensitivity of cardiac conduction system function and homeostasis. *Proc. Natl. Acad. Sci. USA.* 109, E154–E163. doi: 10.1073/pnas.1115165109

Freyermuth, F., Rau, F., Kokunai, Y., Linke, T., Seller, C., Nakamori, M., et al. (2016). Splicing misregulation of SCN5A contributes to cardiac-conduction delay and heart arrhythmia in myotonic dystrophy. *Nat. Commun.* 7:11067. doi: 10.1038/ncomms11067

Fu, J. D., Jiang, P., Rushing, S., Liu, J., Chiamvimonvat, N., and Li, R. A. (2010). Na+/Ca2+ exchanger is a determinant of excitation-contraction coupling in human embryonic stem cell-derived ventricular cardiomyocytes. *Stem Cells* 19, 773–782. doi: 10.1002/scd.2009.0184

Fukuda, T., Hirooka, T., Yamota, K., Nakashima, M., Kawai, T., Yanagi, K., et al. (2020). Exome reports A de novo GNB2 variant associated with global developmental delay, intellectual disability, and dysmorphic...
features. Eur. J. Med. Genet. 63:103804. doi: 10.1016/j.ejmg.2019.103804

Gammage, M. D. (2000). Temporary cardiac pacing. Heart 83, 715–720.

Gao, Z., Rasmussen, T. P., Li, Y., Kutschke, W., Koval, O. M., Wu, Y., et al. (2013). Genetic inhibition of Na+-Ca2+ exchanger current disturbs flight or flight sinoatrial node activity without affecting resting heart rate. Circ. Res. 112, 309–317. doi: 10.1161/circresaha.112.3011903

Gilman, A. G. (1987). G proteins: transducers of receptor-generated signals. Annu. Rev. Biochem. 56, 615–649. doi: 10.1146/annurev.bi.56.070187.003151

Glukhov, A. V., Kalyanasundaram, A., Lou, Q., Hage, L. T., Hansen, B. J., Belevych, A. E., et al. (2010). Functional anatomy of the murine sinus node: high-resolution optical mapping of ankyrin-B heterozygous mice. EMBO J. 29, 4882–4891. doi: 10.1038/emboj.2010.399

Gilmore, R., Demion, M., and Launay, P. (2010). Physiological roles of the TRPM4 channel extracted from background currents. Physiology 25, 155–164. doi: 10.1152/physiol00004.2010

Hagiwara, N., Irisawa, H., and Kameyama, M. (1988). Contribution of two types of calcium currents to the pacemaker potentials of rabbit sino-atrial node cells. J. Physiol. 395, 233–253. doi: 10.1113/jphysiol.1988.sp016196

Han, L., Tang, Y., Li, S., Wu, Y., Chen, X., Wu, Q., et al. (2020). Protective mechanism of SIRT1 on Hcy-induced atrial fibrosis mediated by TRPC3. J. Cell. Mol. Med. 24, 488–510. doi: 10.1111/jcm.14757

Hao, X., Zhang, Y., Zhang, X., Nirmalan, M., Davies, L., Konstantinou, D., et al. (2011). TGFR1-mediated fibrosis and ion channel remodeling are key mechanisms in producing the sinus node dysfunction associated with SCN5A deficiency and aging. Circ. Arrhythm. Electrophysiol. 4, 397–406. doi: 10.1161/circep.110.960807

Harada, M., Luo, X., Qi, X. Y., Tadevosyan, A., Maguy, A., Ordog, B., et al. (2012). Transient receptor potential canonical-3 channel-dependent fibroblast regulation in atrial fibration. Circulation 126, 2051–2064. doi: 10.1161/circulationaha.112.121830

Haron-Khun, S., Weisbrod, D., Bueno, H., Yadim, D., Behar, J., Peretz, A., et al. (2017). SKK (K+) channels are therapeutic targets for the treatment of cardiac arrhythmias. EMBO Mol. Med. 9, 415–429. doi: 10.1002/emmm.201606937

Hasdemir, C., Demion, M., Lazzara, R., and Reynolds, D. W. (2003). G protein-gated inwardly rectifying K+ channel is important for channel activation. Evidence that direct binding of G beta gamma to the GIRK1 G protein-gated inwardly rectifying K+ channel is important for channel activation. Neuron 15, 1133–1143. doi: 10.1016/0896-6273(95)90101-9

Huang, R. T., Xue, S., Xu, Y., Zhou, M., and Yang, Y. Q. (2013). A novel NKX2.5 loss-of-function mutation responsible for familial atrial fibrillation. Int. J. Mol. Med. 31, 1119–1126. doi: 10.3892/ijmm.2013.1316

Huc, S., Monteil, A., Bidaud, I., Barbara, G., Chemin, J., and Lory, P. (2009). Regulation of T-type calcium channels: signalling pathways and functional implications. Biochim. Biophys. Acta. 1793, 947–952. doi: 10.1016/j.bbamcr.2008.11.003

Ishikawa, T., Ohno, S., Murakami, T., Yoshida, K., Mishima, H., Fukuko, T., et al. (2017). Sick sinus syndrome with HCN4 mutations shows early onset and frequent association with atrial fibration and left ventricular noncompaction. Heart Rhythm 14, 717–724. doi: 10.1016/j.hrthm.2017.01.020

James, T. N. (1963). The connecting pathways between the sinus node and A-V node and between the right and the left atrium in the human heart. Am. Heart J. 66, 498–508. doi: 10.1016/S0002-9378(63)90382-X

James, T. N. (1977). The sinus node. Am. J. Cardiol. 40, 965–986. doi: 10.1016/0002-9149(77)90148-0

James, T. N., Sherf, L., Fine, G., and Morales, A. R. (1966). Comparative ultrastructure of the sinus node in man and dog. Circulation 34, 139–163. doi: 10.1161/01.cir.34.1.139

James, T. N., St Martin, E., Willis, P. W. III, and Lohr, T. O. (1996). Apoptosis as a possible cause of gradual development of complete heart block and fatal arrhythmias associated with absence of the AV node, sinus node, and internodal pathways. Circulation 93, 1424–1438. doi: 10.1161/01.cir.93.7.1424

Jeevaratnam, K., Zhang, Y., Guzadhur, L., Duehmke, R. M., Grace, A. A., et al. (2010). Differences in sino-atrial and atrio-ventricular function with age and sex attributable to the Scn5a+/- mutation in a murine cardiac model. Acta Physiol. 200, 23–33. doi: 10.1111/j.1748-1716.2010.02110.x

Ji, S., John, S. A., Lu, Y., and Weiss, J. N. (1998). Mechanosensitivity of the cardiac 2-adrenergic receptor. J. Biol. Chem. 273, 1324–1328. doi: 10.1074/jbc.273.3.1324

Jing, H. L., and Hu, B. J. (1997). Sudden death caused by stricture of the sinus node artery. Am. J. Forensic Med. Pathol. 18, 360–362. doi: 10.1097/00000433-199712000-00009

Ju, Y. K., Chu, Y., Chaulet, H., Lai, D., Gervasio, O. L., Graham, R. M., et al. (2007). Coding and non-coding variants in the SHOX2 gene in patients with early-onset atrial fibrillation. Basic Res. Cardiol. 102,389.

Ju, Y. K., Lee, B. H., Trajanovska, S., Hao, G., Allen, D. G., Lei, M., et al. (2015). The involvement of TRPC3 channels in sinoatrial arrhythmias. Front. Physiol. 6:86. doi: 10.3389/physiol.2015.00086

Kalmus, D., Gerasimenko, O. V., Kalyanasundaram, A., Lou, Q., Hage, L. T., Hansen, B. J., Belevych, A. E., et al. (2010). Functional anatomy of the murine sinus node: high-resolution optical mapping of ankyrin-B heterozygous mice. Eur. J. Med. Genet. 53:103804. doi: 10.1016/j.ejmg.2019.103804
Kalyanasundaram, A., Li, N., Hansen, B. J., Zhao, J., and Fedorov, V. V. (2019). Canine and human sinoatrial node: differences and similarities in the structure, function, molecular profiles, and arrhythmia. J. Vet. Cardiol. 22, 2–19. doi: 10.1016/j.jvcc.2018.10.004

Keith, A., and Flack, M. (1907). The form and nature of the muscular connections between the primary divisions of the vertebrate heart. J. Anat. Physiol. 41(Pt 3), 172–189.

Kennedy, A., Finlay, D. D., Guldenring, D., Bond, R., Moran, K., and McLaughlin, J. (2016). The cardiac conduction system: generation and conduction of the cardiac impulse. Crit. Care Nurs. Clin. North Am. 28, 269–279. doi: 10.1016/j.ccrn.2016.04.001

Kennedy, M. E., Necmy, J., Corey, S., Wickman, K., and Clapham, D. E. (1999). GIRK4 confers appropriate processing and cell surface localization to G-protein-gated potassium channels. J. Biol. Chem. 274, 2571–2582. doi: 10.1074/jbc.274.2.5271

Kertesz, N. J., Towbin, J. A., Clunie, S., Fenrich, A. L., Friedman, R. A., Kearney, K. W., Hong, K. W., Go, M. J., Kim, S. S., Tabara, Y., Kita, Y., et al. (2014). Kokubo, Y., Inamoto, N., Tomoike, H., Kamide, K., Takiuchi, S., Kawano, Y., et al. (2014). Kuratomi, S., Ohmori, Y., Ito, M., Shimazaki, K., Muramatsu, S., Mizukami, K., Kondo, S., Osanai, H., Sakamoto, Y., Uno, H., Tagahara, K., Hosono, H., et al. (2014). Kokunai, Y., Nakata, T., Furuta, M., Sakata, S., Kimura, H., Aiba, T., et al. (2014).

Launay, P., Fleig, A., Perraud, A. L., Scharenberg, A. M., Penner, R., and Kinet, J. P. (2002). TRPM4 is a Ca2+-activated nonselective cation channel mediating cell membrane depolarization. Cell 109, 397–407. doi: 10.1016/s0092-8674(02)00719-5

Leenhardt, A., Lucet, V., Denjoy, I., Grau, F., Ngoc, D. D., and Counel, P. (1995). Catecholaminergic polymorphic ventricular tachycardia in children: a 7-year follow-up of 21 patients. Circulation 91, 1512–1519.

Lei, M., Goddard, C., Liu, J., Leoni, A. L., Royer, A., Fung, S. S., et al. (2005). Sinus node dysfunction following targeted disruption of the murine cardiac sodium channel gene Scn5a. J. Physiol. 567(Pt 2), 387–400. doi: 10.1113/jphysiol.2005.083188

Lesage, F., and Lazdunski, M. (1998). Mapping of human potassium channel genes TREC-1 (KCNK2) and TASK (KCNK3) to chromosomes 1q41 and 2p23. Genomics 51, 478–479. doi: 10.1006/geno.1998.5397

Li, N., Csepe, T. A., Hansen, B. J., Sul, L. V., Kalyanasundaram, A., Zakharkin, S. O., et al. (2016). Adenosine-induced atrial fibrillation: localized reentrant drivers in lateral right atria due to heterogeneous expression of adenosine A1 receptors and GIRK4 subunits in the human heart. Circulation 134, 486–498. doi: 10.1161/circulationaha.115.021165

Li, N., Hansen, B. J., Csepe, T. A., Zhao, J., Ignozzi, A. J., Sul, L. V., et al. (2017). Redundant and diverse intranodal pacemakers and conduction pathways protect the human sinoatrial node from failure. Sci. Transl. Med. 9:eaaem5607. doi: 10.1126/scitranslmed.aam5607

Li, N., Wang, Z.-S., Wang, X.-H., Xu, Y.-J., Qiao, Q., Li, X.-M., et al. (2018). A SHOX2 loss-of-function mutation underlying familial atrial fibrillation. Int. J. Med. Sci. 15, 1364–1372.

Li, Y. D., Hong, Y. F., Zhang, Y., Zhou, X. H., Ji, Y. T., Li, H. L., et al. (2014). Association between reversal in the expression of hyperpolarization-activated cyclic nucleotide-gated (HCN) channel and age-related atrial fibrillation. Med. Sci. Monit. 20, 2292–2297. doi: 10.12659/msm.892305

Liao, J., Li, H., Zeng, W., Sauer, D. B., Belmares, R., and Jiang, Y. (2012). Structural insight into the ion-exchange mechanism of the sodium/calcium exchanger. Science 335, 686–690. doi: 10.1126/science.1215759

Liao, Z., Lockhead, D., Larson, E. D., and Proenza, C. (2010). Phosphorylation and modulation of hyperpolarization-activated HCN4 channels by protein kinase A in the mouse sinoatrial node. J. Gen. Physiol. 136, 247–258. doi: 10.1085/jgp.201010488

Liaqat, K., Schrauw, I., Raza, S. I., Lee, K., Hussain, S., Chakchouk, L., et al. (2018). Identification of CACNA1D variants associated with sinoatrial node dysfunction and deafness in additional Pakistani families reveals a clinical significance. J. Hum. Genet. 64, 153–160. doi: 10.1038/s41008-018-0542-8

Lin, X. F., Luo, J. W., Liu, G., Zhou, Y. B., Jin, Z., and Lin, X. (2018). Genetic mutation of familial dilated cardiomyopathy based on next-generation semiconductor sequencing. Mol. Med. Rep. 18, 4271–4280. doi: 10.3892/mmr.2018.9455

Ling, C. A., and Crouch, M. A. (1998). Theophylline for chronic symptomatic bradycardia in the elderly. Ann. Pharmacother. 32, 837–839. doi: 10.1345/aph.17462

Liu, J., Xin, L., Benson, V. L., Allen, D. G., and Ju, Y. K. (2015). Store-operated calcium entry and the localization of STIM1 and Orai1 proteins in isolated mouse sinoatrial node cells. Front. Physiol. 28, 269–279. doi: 10.1016/j.jphysiol.2015.09.0368

Luo, H., et al. (2003). Lamin A/C truncation in dilated cardiomyopathy with conduction disease. BMC Med. Genet. 4:4. doi: 10.1186/1471-2350-4-4

MacLeod, H. M., Culley, M. R., Huber, J. M., and McNally, E. M. (2003). Lamin A/C truncation in dilated cardiomyopathy with conduction disease. BMC Med. Genet. 4:4. doi: 10.1186/1471-2350-4-4

Mol. Med. Rep. 18, 4271–4280. doi: 10.3892/mmr.2018.9455
Saito, D., Matsubara, K., Yamanari, H., Obayashi, N., Uchida, S., Maekawa, K., et al. (1993). Effects of oral theophylline on sick sinus syndrome. J. Am. Coll. Cardiol. 21, 1199–1204. doi: 10.1016/0735-1097(93)90246-w
Sanchez-Queintana, D., Cabrera, J., Farre, J., Climent, V., Anderson, R., and Ho, S. Y. (2005). Sinus node revisited in the era of electroanatomical mapping and catheter ablation. Heart 91, 189–194.
Sanders, P., Kistler, P. M., Morton, J. B., Spence, S. J., and Kalman, J. M. (2004). Remodeling of sinus node function in patients with congestive heart failure: reduction in sinus node reserve. Circulation 110, 897–903. doi: 10.1161/01.cir.0000139336.69955.ab
Schmidt, C., Wiedmann, F., Tristram, F., Anand, P., Wenzel, W., Lugnbeiil, P., et al. (2014). Cardiac expression and atrial fibrillation-associated remodeling of K<sub>P</sub>2.1 (TREK-1) K<sup>+</sup> channels in a porcine model. Life Sci. 97, 107–115. doi: 10.1016/j.lfs.2013.12.006
Schulze-Bahr, E., Neu, A., Friederich, P., Kaupp, U. B., Breitherd, G., Pongs, O., et al. (2003). Pacemaker channel dysfunction in a patient with sinus node disease. J. Clin. Invest. 113, 1537–1545. doi: 10.1172/jci16387
Schweitzer, P., and Mark, H. (1980). The effect of atropine on cardiac arrhythmias and conduction. Part 2. Am. J. Heart 100, 255–261. doi: 10.1001/0000139336.6955.ab
Schweizer, P. A., Duhme, N., Thomas, D., Becker, R., Zehelein, J., Draguhn, A., et al. (2014). The symptom complex of familial sinus node dysfunction and disease. J. Clin. Invest. 128, 524–535. doi: 10.1172/JCI71835
Sherraishi, I., Takamatsu, T., Minamikawa, T., Onouchi, Z., and Fujita, S. (1992). Calsequestrin 2 (CASQ2) mutations increase expression of calreticulin and ryanodine receptors, causing catecholaminergic polymorphic ventricular tachycardia. J. Clin. Invest. 117, 1814–1823. doi: 10.1172/jci131080
Sonoura, T., Koderu, S., Shaky, S., and Kanda, J. (2019). Efficacy of clobalost for sick sinus syndrome to avoid permanent pacemaker implantation: a retrospective case-control study. J. Cardiol. 74, 328–332. doi: 10.1016/j.jcc.2019.03.007
Squire, J. M. (1997). Architecture and function in the muscle sarcomere. Curr. Opin. Struct. Biol. 7, 247–257. doi: 10.1016/s959-440x(97)80033-4
Stallmeyer, B., Kuft, J., Kotthoff, S., Zumhagen, S., Vowinkel, K., Rinné, S., et al. (2017). A mutation in the G-protein gene GB2 causes familial sinus node and atrioventricular conduction dysfunction. Circ. Res. 120, e33–e64. doi: 10.1161/ Circresaha.116.310112
Stallmeyer, B., Zumhagen, S., Denjoy, I., Duthoit, G., Hébert, J. L., Ferrer, X., et al. (2012). Mutational spectrum in the Ca(2+)–activated cation channel gene TRPM4 in patients with cardiac conductance disturbances. Hum. Mutat. 33, 109–117. doi: 10.1002/humu.21599
Stieber, I., Hofmann, F., and Ludwig, A. (2004). Pacemaker channels and sinus node arrhythmia. Trends Cardiovasc. Med. 14, 23–28. doi: 10.1016/j.tcm.2003.09.006
Sumitomo, N., Sakurada, H., Taniguchi, K., Matsumura, M., Abe, O., Miyashita, M., et al. (2007). Association of atrial arrhythmia and sinus node dysfunction in patients with catecholaminergic polymorphic ventricular tachycardia. Circ. J. 71, 1606–1609.
Takahashi, K., Cecchin, F., Fortescue, E., Berul, C. I., Alexander, M. E., Walsh, E. P., et al. (2009). Permanent atrial pacing lead implant route after Fontan operation. Pacing Clin. Electrophysiol. 32, 779–785. doi: 10.11151.1540-8159.2009.02365.x
Tan, B. H., Iturralde-Torres, P., Medeiros-Domingo, A., Nava, S., Tester, D. J., Valdivia, C. R., et al. (2007). A novel C-terminal truncation SCN5A mutation from a patient with sick sinus syndrome, conduction disorder and ventricular tachycardia. Circ. Res. 76, 409–417. doi: 10.1016/j.circres.2007.08.006
Thery, C., Gosselin, B., Lekkrief, J., and Warembourg, H. (1977). Pathology of sinoatrial node. Correlations with electrocardiographic findings in 111 patients. Am. J. Heart 93, 735–740. doi: 10.1016/s0002-8703(77)80670-7
Torrence, A. G., Mesirca, P., Neco, P., Rizzetto, R., Dubel, S., Barrere, C., et al. (2016). L-type Cav3.1 channels regulate ryanodine receptor-dependent Ca2+ release during sino-atrial node pacemaker activity. Cardiovasc. Res. 109, 451–461. doi: 10.1093/ercr/cvw006
Torrence, A. G., Zhang, R., Wang, H., Zaini, A., Kim, B., Yue, X., et al. (2017). Contribution of small conductance K<sup>+</sup> channels to sinoatrial node pacemaker activity: insights from atrial-specific Naf<sup>+</sup> /Ca<sup>2+</sup> exchange knockout mice. J. Physiol. 595, 3848–3865. doi: 10.1113/jp274249
Torrence, A. G., Zhang, R., Zaini, A., Giani, J. F., Kang, J., Lamp, S. T., et al. (2015). Burst pacemaker activity of the sinoatrial node in sodium-calcium exchanger knockout mice. Proc. Natl. Acad. Sci. U.S.A. 112, 9769–9774. doi: 10.1073/pnas.1505670112
Tracy, C. M., Epstein, A. E., Darbar, D., DiMarco, J. P., Dunbar, S. B., Estes, N. M., et al. (2013). 2012 ACCF/AHA/HRS focused update incorporated into the ACCF/AHA/HRS 2008 guidelines for device-based therapy of cardiac rhythm abnormalities: a report of the American College of Cardiology Foundation/American Heart Association Task Force on Practice Guidelines and the Heart Rhythm Society. J. Am. Coll. Cardiol. 61, e6–e75.
Ueda, K., Nakamura, K., Hayashi, T., Inagaki, N., Takahashi, M., Arimura, T., et al. (2004). Functional characterization of a trafficking-defective HCN4 mutation, D553N, associated with cardiac arrhythmia. J. Biol. Chem. 279, 27194–27198. doi: 10.1074/jbc.M411953200
Unudruthi, S. D., Wu, X. Q., Qian, L., Amari, F., Onal, B., Li, N., et al. (2016). Two-pore K<sup>+</sup> channel TREK-1 regulates sinoatrial node membrane excitability. J. Am. Heart Assoc. 5:e002865. doi: 10.1161/jaha.115.002865
van den Berg, M. P., Wilde, A. A., Viersma, T. J. W., Brouwer, J., Haakasma, J., van der Hout, A. H., et al. (2001). Possible bradycardic mode of death and successful pacemaker treatment in a large family with features of long QT syndrome type 3 and Brugada syndrome. J. Cardiovasc. Electrophysiol. 12, 630–636. doi: 10.1046/j.1540-8197.2001.00630.x
van Elf, V. W. W., Stefanovic, S., van Duijvenboden, B., Bakker, M., Wakker, V., de Gier-de Vries, C., et al. (2019). Transcriptome analysis of mouse and human sinoatrial node cells reveals a conserved genetic program. Development 146:dev173161. doi: 10.1242/dev.173161
Vavetsi, S., Nikolou, N., Tsarouhas, K., Lymeropoulos, G., Kouzanidis, I., Kafantaris, L., et al. (2008). Consecutive administration of atropine and isoproterenol for the evaluation of asymptomatic sinus bradycardia. *European Heart Journal*, 110, 1176–1181. doi: 10.1093/eurheartj/eun211

Verhaeghe, L., and van der Haute, L. (1967). Arterial blood supply of the human sinus node. *Br. Heart J.* 29, 801–806. doi: 10.1136/brhtj.29.5.801

Wang, F., Liu, J., Hong, L., Liang, B., Graff, C., Yang, Y., et al. (2013). The phenotype characteristics of type 13 long QT syndrome with mutation in KCNJ5 (Kir3.4-G387R). *Heart Rhythm* 10, 1500–1506. doi: 10.1016/j.hrthm.2013.07.022

Wasada, T., Katsumorii, K., Hasumi, S., Kasanuki, H., Arai, H., Saeki, A., et al. (1995). Association of sick sinus syndrome with hyperinsulinemia and insulin resistance in patients with non-insulin-dependent diabetes mellitus: report of four cases. *Intern. Med.* 34, 1174–1177. doi:10.2169/internalmedicine.34.1174

Weeke, P., Purvez, B., Blais, M., Shortt, L., Ingram, C., Cucuera, G., et al. (2014). Candidate gene approach to identifying rare genetic variants associated with lone atrial fibrillation. *Heart Rhythm* 11, 46–52. doi: 10.1016/j.hrthm.2013.10.025

Wessels, A., and Sedmara, D. (2003). Developmental anatomy of the heart: a tale of mice and man. *Physiol. Genomics* 15, 165–176. doi: 10.1152/physiogenomics.00033.2003

Wickman, K., Nemec, J., Gendler, S. J., and Clapham, D. E. (1997). Partial inactivation of the mouse Kir4.4 channels. *Genomics* 40, 395–401. doi: 10.1006/geno.1997.4599

Wilde, A. A. M., and Amin, A. S. (2018). Clinical spectrum of SCN5A mutations: long QT syndrome, Brugada Syndrome, and cardiomyopathy. *JACC Clin. Electrophysiol.* 4, 569–579. doi: 10.1016/j.jcpace.2018.03.006

Wilders, R. (2018). Cellular mechanisms of sinus node dysfunction in carriers of the SCN5A-E161K mutation and role of the H558R polymorphism. *Front. Physiol.* 9:1795. doi: 10.3389/fphy.2018.01795

Wolf, C. M., and Berul, C. I. (2006). Inherited conduction system abnormalities—one group of diseases, many genes. *J. Cardiovasc. Electrophysiol.* 17, 446–455. doi:10.1111/j.1540-8167.2006.00427.x

Wolf, R. M., Glyn, P., Hashemi, S., Zarei, K., Mitchell, C. C., Anderson, M. E., et al. (2013). Atrial fibrillation and sinus node dysfunction in human ankyrin-B syndrome: a computational analysis. *Am. J. Physiol. Heart Circ. Physiol.* 304, H1253–H1266. doi: 10.1152/ajpheart.00734.2012

Wolf, R. M., Mitchell, C. C., Christensen, M. D., Mohler, P. J., and Hund, T. J. (2010). Defining new insight into atypical arrhythmia: a computational model of ankyrin-B syndrome. *Am. J. Physiol. Heart Circ. Physiol.* 299, H1505–H1514. doi: 10.1152/ajpheart.00503.2010

Woo, G. W., Schofield, R. S., Pauly, D. F., Hill, A. J., Conti, J. B., Kron, J., et al. (2008). Incidence, predictors, and outcomes of cardiac pacing after cardiac transplantation: an 11-year retrospective analysis. *Transplantation* 85, 1216–1218. doi:10.1097/TP.0b1013e3181667677

Woods, W. T., Uthraler, F., and James, T. N. (1976). Spontaneous action potential of cells in the canine sinus node. *Circ. Res.* 39, 76–82.

World Health Organization (2020). WHO Director-General’s Opening Remarks at the Media Briefing on COVID-19 11 March 2020. Geneva: World Health Organization.

Wu, X., Zagranichnaya, T. K., Gurda, G. T., Eves, E. M., and Villareal, M. L. (2004). A TRPC1/TRPC3-mediated increase in store-operated calcium entry is associated with heart disease in carriers of the CACNA1C D604S mutation. *Transplantation* 8457–8465. doi: 10.3892/mmr.2018.8873

Xia, J., Zhang, J., Zhao, X., Li, H., Gao, Y., et al. (2013). Atrial autonomic denervation for the treatment of long-standing symptomatic sinus bradycardia in non-elderly patients. *J. Interv. Card. Electrophysiol.* 43, 151–159.

Zhu, Y., Feng, Z., Cheng, W., and Xiao, Y. (2018). MicroRNA-34a mediates atrial fibrillation through regulation of Ankyrin-B expression. *Mol. Med. Rep.* 17, 8457–8465. doi: 10.3892/mmr.2018.8873

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The remaining authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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