Loss-of-Function SCN5A Mutations Associated With Sinus Node Dysfunction, Atrial Arrhythmias, and Poor Pacemaker Capture

Running title: Chiang et al.; SCN5A mutations and poor pacemaker capture

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Abstract:

**Background** - Cardiac device implantation can be complicated by inability to adequately place leads due to significant lead capture issues. This study sought to determine if there are genetic bases that underlie poor lead capture.

**Methods and Results** - Retrospective review of all patients with structurally normal hearts who underwent new device implantation at Texas Children’s Hospital between 2009-2014 was performed. Patients with inability to capture at 10V or a final capture threshold ≥3V at 0.4ms during implant were analyzed. Among a total of 136 patients (median age 13 years, range 3 days-46 years), 11 patients (8.1%) who underwent dual chamber device implantation had elevated thresholds in the atria (4), ventricle (3), or both chambers (4) (atrial-lead threshold: 4.7±4.3V vs. 0.7±0.3V; ventricular-lead: 3.0±3.3V vs. 0.7±0.3V). All 11 patients presented with sinus node dysfunction and 10 had atrial arrhythmias. At implant, inability to find atrial capture was seen in 4 patients. Three demonstrated intermittent complete loss of ventricular capture after implantation: 1 has recurrent syncope, 2 eventually died. Genetic testing performed in 10 demonstrated 7 patients with 6 distinct SCN5A mutations, all predicted to be severe loss-of-function mutations by bioinformatic analyses. In the remaining patients, although putative pathogenic mutations were not found, multiple SCN5A polymorphisms were identified in 2 and a desmin mutation in 1.

**Conclusions** - This study suggests that significant capture issues at implant may be due to loss-of-function SCN5A mutations, providing new insights into SCN5A function. Recognition of this association may be critical for planning device implantation strategies and patient follow-up.

**Key words:** SCN5A, mutation, pacemaker, atrial tachyarrhythmia, sinus node dysfunction
Introduction

Significant issues with pacemaker or implantable cardioverter defibrillator (ICD) lead placement can occur during device implantation due to an inability to obtain a reasonable lead capture threshold. Although infrequent, this complication has led to difficulty or even inability to place leads, resulting in compromises to patient care and potentially adverse outcomes. We noted that in some of these patients, capture thresholds undulated over time, leading us to wonder whether these alterations might be secondary to ion channel dysfunction. A previous study associated heterozygous mutations in \textit{SCN5A}, which codes for the cardiac voltage-gated sodium channel (Na\textsubscript{v}1.5), with sick sinus syndrome and atrial inexcitability but never elaborated upon capture issues.\textsuperscript{1} Later, our center was the first to report a patient with atrial arrhythmias found to have poor capture thresholds during lead placement.\textsuperscript{2} Genetic testing revealed that she harbored a homozygous loss-of-function mutation in \textit{SCN5A}. With this in mind, the present study reviewed all patients with poor pacemaker capture at Texas Children’s Hospital from 2009 to 2014 as well as all patients with known \textit{SCN5A} mutations at our center in order to test the hypothesis that loss-of-function \textit{SCN5A} mutations are associated with poor pacemaker capture.

Methods

A retrospective review of all patients at Texas Children’s Hospital (Houston, Texas) who underwent new pacemaker or ICD implantation between 2009 and 2014 was performed with approval from the Internal Review Board. Dates were chosen in the current era where genetic testing was available and would have been performed at our institution. Patients with congenital heart disease and heart transplant were excluded. Patients with cardiomyopathy, including left ventricular noncompaction (LVNC) were included. Records were reviewed for significant difficulties with lead capture at the time of implant as reported by the implanting physician. Any
patient with documented capture thresholds >10V or a final capture threshold ≥3V at a pulse width of 0.4-0.5ms was included. Final capture thresholds reported in this study reflect the final threshold at implant. If multiple sites were attempted with inability to capture at 10V and the final implant site had a capture threshold of 1.5V, the final capture threshold was recorded as 1.5V for the purposes of this study. If a lead was ultimately not placed due to inability to capture at 10V, the threshold was recorded as 10V. Lead sensing at implant was also recorded, although this was not the primary focus. Leads with sensing <0.3mV were considered abnormal. Patients with lead sensing problems without capture issues were excluded. In addition, a retrospective review of all other patients with SCN5A mutations at Texas Children’s Hospital was performed.

Data collection included demographics, clinical history, review of electrocardiograms, genetic testing results and device data including implant records.

**Genetic Testing**

After informed consent was obtained, the patients’ blood sample was sent to either Familion (New Haven, Connecticut) or John Welsh Cardiovascular Diagnostic Laboratory (Houston, Texas). For Familion, the genomic DNA isolated from blood was PCR amplified and used for direct sequencing of the complete open reading frame, splice junctions, and flanking regions of the genes of interest. For John Welsh Laboratory, the isolated genomic DNA was used for PCR amplification of the exons and splice junctions of the genes of interest before direct sequencing. Of the 10 patients with genetic testing, 2 were tested by Familion using their long QT syndrome panel which included KCNQ1, KCNH2, SCN5A, KCNE1, and KCNE2 while 8 were tested by the John Welsh Laboratory. Of these 8 patients, 5 were tested using the AV block panel which included LMNA, DES, EMD, NKX2.5, SCN5A, and SCN1B; 1 was tested using the Brugada Syndrome/J-Wave Syndrome panel which included CACNA1C, CACNB2, GPD1L, KCND3,
KCNE3, KCNJ8, HCN4, SCN1B, SCN2B, SCN3B, and SCN5A; and 2 were tested only for SCN5A.

Statistical Analysis

Data are presented as mean ± SD.

Results

Patient Population

During the time period between 2009 and 2014, a total of 425 patients underwent a device-related procedure and 136 of these underwent new device implant (60 pacemaker, 76 ICD) at a median age of 13 years (range 3 days-46 years). New devices were implanted for the following primary reasons: congenital or surgical AV block (43), hypertrophic cardiomyopathy (29), long QT syndrome (27), sinus node dysfunction (15), dilated cardiomyopathy (DCM; 8), Brugada syndrome (BrS; 4), left ventricular noncompaction (LVNC; 2), catecholaminergic polymorphic ventricular tachycardia (2), Becker’s cardiomyopathy (2), arrhythmogenic right ventricular dysplasia (1), and other (3). A total of 200 leads were placed (131 endocardial, 69 epicardial).

Of the 136 patients, 11 had significant capture issues at implant (median age 11 years, range 3 days-17 years) and met study criteria (Figure 1A). Five of the 11 patients were male and 8 were of Hispanic origin. Dual chamber pacemakers were attempted in 9 (6 epicardial, 3 transvenous) and dual chamber ICDs were placed in 2 (1 epicardial, 1 transvenous). At implantation, inadequate capture was seen in the atrium in 4 patients, ventricle in 3 patients, and in both atrium and ventricle in 4 patients (Table 1). Both mean atrial (4.7±4.3V vs. 0.7±0.3V) and ventricular thresholds (3.0±3.3V vs. 0.7±0.3V) at implant were higher compared to those without reported capture issues. In addition, 8 of 11 (73%) patients had extremely low or essentially no sensing at implant (Table 1). Median follow-up from implant for the 11 patients was 16 months (range 15
days-5 years). There were 2 deaths at 15 days and 11 months after device implant.

**Genetic Analyses**

Genetic testing was performed in 10 of the 11 patients with capture issues; results were not available prior to device implantation in any of the patients. Among the 10 patients with genetic testing, 6 distinct Class I SCN5A mutations (including one novel mutation) were found in 7 patients (70%) (Table 1). In 2 patients, in whom no putative mutations were found, multiple single nucleotide polymorphisms (SNPs) in SCN5A were reported (Table 1). In the last patient, no SCN5A mutations or SNPs were identified; however, a variant of unknown significance in DES, which encodes desmin, was seen (Figure 1B). The cohort of 7 patients with putative Class I SCN5A mutations consisted of 4 females, two of whom are identical twins, and 3 males, with age of presentation ranging from newborn to 15 years. Six of the 7 patients were Hispanic. Missense mutations were analyzed in terms of conservation across species (Figure 2) as well as using PolyPhen-2, which predicts the severity of the resultant amino acid substitution (Table 1). 3

**Clinical Features**

Among these 11 patients, one patient did not undergo genetic testing. This patient had sinus node dysfunction and complete heart block with poor sensing and capture in the ventricle necessitating placement of a left ventricular epicardial lead (Figure 1A). In the remaining 10 patients, all of whom were genetically tested, 9 had isolated cardiac issues, while one had multiple medical issues including developmental delay, seizure disorder, prolonged QT (with negative genetic testing), and DCM. Echocardiography demonstrated structurally normal hearts with normal chamber size and normal biventricular function in 9 patients. Two patients (including one with multiple medical problems) demonstrated LVNC (Table 1).

Despite these individual differences, all 11 patients demonstrated sinus node dysfunction,
10 of whom had documented atrial arrhythmias including 4 with atrial flutter, 2 with atrial flutter/atrial fibrillation, and 4 with ectopic atrial tachycardia (Figure 1A). The single patient without atrial arrhythmia was also the one without genetic testing. However, that patient did have multiple atrial high rate episodes at 225 bpm (higher than expected sinus tachycardia rates) at times consistent with sleep although definitive atrial tachycardia was not documented by electrograms. At implant, there was complete quiescence of atrial activity and inability to find atrial capture in 4 patients. In 2 patients, an epicardial atrial (A-)lead could not be placed due to inadequate capture threshold and therefore, an atriotomy was performed and the lead inserted into the endocardium via an epicardial approach through the right atrial appendage. Three patients demonstrated intermittent complete loss of ventricular capture after implantation, 2 of whom eventually died. The remaining patient has a history of undulating capture thresholds with recurrent syncope due to intermittent loss of ventricular capture during times of illness and is currently undergoing evaluation for possible heart transplantation.

**Inclusion Analysis of All Patients with SCN5A Mutations**

Due to the high incidence of SCN5A mutations among patients with poor pacemaker capture (7 out of 10 tested), *all* patients with known SCN5A mutations at Texas Children’s Hospital were reviewed to determine how many patients with known SCN5A mutations also had pacemaker capture issues. There were a total of 27 patients at our institution with 20 distinct SCN5A mutations (large circle in Figure 3A). Of these 27 patients, 15 underwent placement of a pacemaker or ICD. Seven of these 15 patients demonstrated pacemaker capture issues and are included in our cohort of 11 patients. The remaining 8 patients did not exhibited sinus node dysfunction, atrial arrhythmias, or pacemaker capture issues (medium circle in Figure 3A). In other words, 26% of all patients with confirmed SCN5A mutations at our institution had capture
issues (7 out of 27) (small circle in Figure 3A).

Of the 7 patients with SCN5A mutations and pacemaker capture issues, only one had a clinical diagnosis of Brugada syndrome (BrS); however during review of all ECGs for this study, in one other patient, a single ECG (among many performed) demonstrated a Type I Brugada pattern. The other 5 patients demonstrated nonspecific conduction delays and ST changes without definitive Brugada changes. None of these patients underwent provocative drug testing to evaluate for BrS. In contrast, of the 8 patients with SCN5A mutations without capture issues, 3 had a definitive clinical diagnosis of BrS and 5 had a diagnosis of long QT syndrome (Figure 3B). Bioinformatic analysis using PolyPhen-2 of the SCN5A mutations in these patients could not distinguish between patients with and without capture issues.3 The 7 cases with definitive SCN5A mutations and poor capture are described below as short vignettes.

Case 1
A 4-year-old Hispanic female initially presented with sinus node dysfunction. During placement of a dual-chamber epicardial pacemaker, an A-lead could not be placed due to inability to capture despite multiple sites including both right and left atrium at maximal outputs. A single epicardial ventricular (V-)lead with an implant pacing threshold of 8.6V was placed. Her V-threshold improved with time, remaining elevated but stable (2.75V at 0.4ms). Over the ensuing four years, she has been admitted multiple times for atrial flutter requiring direct current (DC) cardioversion. She has remained clinically stable for over one year on metoprolol and warfarin.

Case 2
A newborn Hispanic male was initially diagnosed with ventricular ectopy, atrial-ventricular (AV) block, and LVNC with depressed function. Three days after birth, he underwent placement of an uncomplicated epicardial, dual-chamber pacemaker with normalization of systolic function.
At age two years, he had several episodes of syncope and seizures associated with a viral illness, as well as runs of atrial tachycardia. Pacemaker interrogation showed complete non-capture by the V-lead. During pacemaker revision, an adequate epicardial ventricular pacing site could not be found, until an area was finally located after 2 hours in the left ventricle with a final threshold of 1.4V at 1ms. The existing A-lead was left untouched with a threshold of 1.25V at 0.52ms. A few days later, the ventricular threshold increased to 2.25V at 1ms and has since fluctuated with thresholds >5V at times. The thresholds wax and wane with highest thresholds occurring during times of acute illness. Recently, he had another episode of syncope associated with mild viral symptoms and his V-lead demonstrated complete lack of capture at maximal output. During admission, he demonstrated episodes of nonsustained atrial and ventricular tachycardia (VT) for which he was treated with amiodarone and metoprolol. With resolution of his viral illness, all arrhythmias quiesced and his myocardial capture returned.

Case 3
A 20-month-old Hispanic boy initially presented with an unresponsive episode after developing a fever. Work-up revealed atrial flutter/fibrillation with variable conduction, sinus bradycardia with a junctional escape, and sinus pauses up to 4 seconds. Due to a single ECG noted to demonstrate Type I Brugada changes he was diagnosed with BrS. During epicardial lead placement, poor sensing and capture of both the right atrium and right ventricle was encountered resulting in the need to convert a subxiphoid incision into a full sternotomy. After much trial, the V-lead was placed on the left ventricular apex. The A-lead was secured into the right atrial endocardium through a right atrial appendage incision. Both leads demonstrated only intermittent capture at full 10V output.

Post-operatively, he suffered an acute and extensive thrombotic stroke. Four days later,
the patient had multiple episodes of pacemaker capture loss and hypotension. Despite maximal pacemaker output, in a non-acidotic, non-hypokalemic state, intermittent loss of capture persisted. Due to bradycardia and low cardiac output with multi-organ system failure and evidence of irreversible neurologic injury, support was withdrawn.

**Case 4**

A 15-year-old Hispanic male presented with atrial flutter requiring DC cardioversion. He was subsequently found to have sinus node dysfunction with multiple sinus pauses up to 4.5 seconds. Intracardiac mapping during an electrophysiology study found significantly decreased atrial voltages, inability to capture portions of atrial tissue even at maximum output, and intra-atrial and inter-atrial conduction delays. During subsequent placement of a transvenous dual-chamber pacemaker, the A-lead had to be placed along the low atrial septum to achieve capture. The V-lead was placed without issues.

**Case 5**

As previously reported, a 6-year-old Hispanic female initially presented with sinus bradycardia and frequent sinus pauses but was lost to follow-up for 5 years. At age 11, she presented with recurrent sustained VT after an aborted sudden cardiac death, requiring extracorporeal membrane oxygenation support for severely depressed biventricular function. Because her arrhythmia was refractory to medications, she underwent placement of a pacemaker/defibrillator. Temporary A-leads could not be placed due to inability to capture at full 10V output. Temporary V-leads paced intermittently at full 10V output. The following day, a dual-chamber epicardial ICD was attempted. Once again, despite multiple attempts to place an A-lead, the atria were entirely quiescent and atrial capture could not be achieved. Nevertheless, an epicardial lead was placed in case capture could be later achieved. Similarly for the V-lead, only intermittent ventricular
capture was obtained despite attempts at multiple sites. Over time, the patient’s ventricular function improved and her epicardial system demonstrated significant improvement with her most recent atrial and ventricular capture thresholds respectively 0.75V at 0.6ms and 1.5V at 0.6ms at the latest interrogation. She has demonstrated nonsustained atrial tachycardia on pacemaker interrogation and a single ECG was found to demonstrate a Type I Brugada pattern. This patient has an identical twin sister described in the next case.

**Case 6**

Like her identical twin from Case 6, this patient presented at age 12 with sinus pauses of up to 5 seconds associated with lightheadedness and dizziness. Given her sinus node dysfunction and symptoms as well as her family history, a dual chamber transvenous ICD was implanted for primary prevention. Atrial and ventricular capture thresholds at implant were poor and variable. Pacemaker interrogation has revealed episodes of atrial tachycardia. Subsequent genetic testing confirmed the same A1924T homozygous mutation in SCN5A.  

**Case 7**

A 5-year-old Caucasian female presented with near syncope in sustained atrial tachycardia. She underwent successful DC cardioversion and subsequent Holter monitoring demonstrated episodes of atrial flutter, sinus node dysfunction and sinus pauses. A- and V-threshold shortly after implant of a dual chamber pacemaker were 2V at 0.09ms and 3V at 0.15ms, respectively, and have remained persistently high in the succeeding months although occasionally the thresholds have improved to 1V at 0.15ms and 2V at 0.27ms, respectively.

**Mutation Analyses**

A novel SCN5A mutation was discovered in patient 7 of our cohort and is predicted to result in a truncated Na\textsubscript{v}1.5 (Figure 4). This mutation is a deletion of nucleotide number 100 (100delC) in
exon 2 of SCN5A (counting from the start codon). This leads to a premature translation stop codon corresponding to the 63rd amino acid after the deletion (R34Afs*63), which would lead to a protein product of only 95 amino acids long (Figure 4), with the following sequence:
MANFLLPRGTSSFRRFTRESLAAIEKRMAEKQAAAQPPCRAERGCPRRRLPGPSWTCRPPKSCQISMAIH
PKSSSESPWRTWPSIAPKRLSSY.

Thus, in our cohort, both Cases 1 and 7 harbor mutations that predict a severely truncated Na\textsubscript{v}1.5 which would most likely undergo nonsense mediated decay, leaving the patients with at most 50% functional Na\textsubscript{v}1.5 (Figures 3 and 5; Table 1).\textsuperscript{4} Case 2 harbors a Na\textsubscript{v}1.5 mutation, G1743R, which is predicted to completely abolish its trafficking to the cell membrane based on \textit{in vitro} studies in HEK293 cells.\textsuperscript{5} The same study also reports that this trafficking defect may be partially rescued with mexiletine or quinidine treatment, although the underlying mechanism was not explored.\textsuperscript{5} Therefore, Case 2 is predicted to have only 50% of functional Na\textsubscript{v}1.5, similar to Cases 1 and 7 (Table 1).

Case 3 harbors the R1512W mutation which based on an earlier \textit{in vitro} study demonstrated a left-shift in both activation and inactivation curves at room temperature.\textsuperscript{6} While that study argues that the R1512W mutation would lead to an overall increase in \(I_{Na}\) against conventional wisdom, a more likely explanation (as conceded by the authors in their discussion) is that because of the left shift, some Na\textsubscript{v}1.5 channels are already activated and then inactivated during resting membrane potential. Therefore, when the cell is depolarized, these channels will not be available to fire, leading to a decreased \(I_{Na}\) and blunting of the action potential upstroke. Furthermore, this left shift may be even more pronounced at body temperature (37°C) since this study was conducted at room temperature (22°C).\textsuperscript{6} In other words, the R1512W mutation may also severely compromise Na\textsubscript{v}1.5 function although further studies, especially \textit{in vivo}, are
needed to rectify the current controversial findings.

The D1275N mutation in Case 4 was initially reported in a Dutch family who presented with atrial standstill but was subsequently associated with a range of cardiac diseases including DCM, sinus node dysfunction, atrial and ventricular tachyarrhythmias, conduction disease, and ventricular enlargement without impaired contractility.\textsuperscript{7-10} However, a number of \textit{in vitro} studies that compared wild-type to D1275N channels in heterologous expression systems failed to demonstrate major differences in the biophysical characteristics of the D1275N mutant channel.\textsuperscript{7, 11} In order to resolve these apparently contradictory findings, a recent study generated a mouse model expressing the human D1275N mutant channel and showed that not only did the animals develop slowed conduction, heart block, atrial fibrillation, VT, and a DCM phenotype, but cardiomyocytes isolated from these animals showed a significantly depressed $I_{\text{Na}}$, when compared to mice expressing the wild-type human channel.\textsuperscript{12} This study underscores the importance of not over-interpreting \textit{in vitro} studies and the need to use multiple model systems especially higher organisms, when trying to understand how a particular mutation may affect a patient \textit{in vivo}. With this study in mind, the D1275N mutation in Case 4 most likely causes a severe decrease in Na\textsubscript{V}1.5, perhaps not dissimilar in magnitude to the mutations in Cases 1 to 3 (Table 1).

The final mutation, A1924T, found in the twins of Cases 5 and 6, was previously characterized \textit{in vitro}. When expressed in \textit{Xenopus} oocytes, the A1924T Na\textsubscript{V}1.5 mutant showed a 9mV negative voltage shift of the steady-state activation curve at room temperature.\textsuperscript{6} This could potential lead to a decrease in $I_{\text{Na}}$ since at resting membrane potential a greater fraction of Na\textsubscript{V}1.5 would already be opened and therefore unavailable when the myocyte is actually depolarized. Given that our patients are homozygous for this mutation, this could lead to an even
greater decrease in total $I_{Na}$ during phase 0 of the cardiac action potential, and thereby a more severe phenotype.

Discussion

In this study, we identified a cohort of patients with poor pacemaker capture at implant. We found that all demonstrated sinus node dysfunction and nearly all of them had clinically significant atrial arrhythmias (Figure 1 and Table 1). A majority of these patients demonstrated loss-of-function SCN5A mutations or multiple SCN5A polymorphisms (Figure 5 and Table 1).

Among known mutations of key ion channel genes, mutations in SCN5A are unique in that they have been implicated in a multitude of cardiac diseases including ones with dissimilar electrophysiological profiles.\textsuperscript{13} For examples, gain-of-function SCN5A mutations may cause long QT syndrome type 3 while loss-of-function mutations may cause conduction disease, BrS, and sick sinus syndrome.\textsuperscript{1, 13, 14} To complicate matters further, the same SCN5A mutation may also present differently even within the same family, for example with features of long QT syndrome, BrS, and conduction disease, in a phenomenon now referred to as “overlap syndromes”.\textsuperscript{15}

Finally, common “benign” variants or SNPs can also potentially modulate phenotype severity by affecting both the biophysical properties of $\text{Na}_{1.5}$ or its expression level.\textsuperscript{13}

A secondary review of all SCN5A patients in our institution did not demonstrate poor pacemaker capture among patients with definitive clinical diagnosis of BrS or long QT syndrome (Figure 3). These findings raise the question whether patients with sinus node dysfunction, atrial arrhythmias and poor pacemaker capture represent a unique clinical phenotype, early BrS, or whether they are part of this continuum in the spectrum SCN5A overlap syndromes.\textsuperscript{13, 15}

In our cohort, pacemaker capture difficulties were seen in both the atrium and ventricle. While we acknowledge the possibility that atrial capture issues may not be recognized in patients
with SCN5A mutations who undergo a single chamber ICD, interestingly there have been no reported cases of inability to capture the ventricle among patients with SCN5A mutations. Similarly, Conte et al. reported that 7.5% of their children with drug-induced Brugada changes had sinus node dysfunction; however they did not comment on whether or not their patients had any complications during ICD implant.16

A clinically important observation from the present study is that the pacemaker capture threshold seems to be highest during acute episodes of illness, which is often at the time of implantation. However, once the patients recover, capture thresholds tend to improve but may fluctuate over time. Although the mechanism by which fever elicits Brugada ECG changes is not clear, fever and elevated temperatures are thought to inactivate NaV1.5 channels.15 Acute increases in metabolic demand secondary to illness may result in alterations within the cardiac milieu such that cardiomyocytes require full functioning of all available NaV1.5 channels in order to remain excitable and conductive. Since our patients have a decreased reserve of functional NaV1.5 channels secondary to their genetic mutations, cardiomyocyte (especially atrial myocyte) NaV1.5 channel dysfunction may become prominent in the setting of fever, acute stress or metabolic demand. At implant, capture thresholds are tested by pacing the heart rates above baseline sinus rhythm. We speculate whether pacing at heart rates above baseline may result in the inability to depolarize a critical threshold of atrial myocytes and thus result in loss of pacemaker capture. Whether or not pacing at slower heart rates during implant would result in pacemaker capture has not been tested.

This study also underscores the importance of planning prior to pacemaker implantation in this patient population. If capture thresholds are unacceptably high, we recommend making all efforts to try several different pacing sites. As in many of our cases, we were eventually able to
achieve a successful site for pacing, even though this may have required multiple attempts. Ultimately if no adequate location is found, we suggest lead implantation be considered even if pacing thresholds are unacceptable, since thresholds may subsequently normalize once the patient improves clinically (as illustrated by Case 5). A dilemma arises when considering the possibility of complete loss of ventricular capture as was seen in 3 of the patients, which were not secondary to significant hyperkalemia and/or acidosis based on clinical history. In Case 3, inability to capture and low cardiac output contributed to multi-system organ failure. In the surviving patient (Case 2), syncope and seizures as a result of complete loss of ventricular capture has resulted in consideration for possible heart transplant. At this time he is being managed by weekly threshold interrogation and inpatient admission for any illness. The remaining patient with intermittent loss of ventricular capture had multiple medical problems including DCM. A hospital admission for respiratory distress and seizures was notable for VT and bradycardia with heart rates dropping to the 30’s suggesting loss of ventricular capture (lower rate limit of pacemaker was 70bpm). Unfortunately, documentation of bradycardia was not available to confirm loss of pacemaker capture. Interrogation of the pacemaker the following day demonstrated appropriate capture thresholds. She was ultimately discharged home. Four days later she died in her sleep. Although we cannot confirm the cause of death, this case highlights the fact that at minimum, close follow-up and monitoring may be necessary in these cases, particularly during times of acute illness. If the clinical phenotype becomes more malignant and unpredictable, more aggressive approaches, including consideration for transplant, may need to be considered.

Limitations

This is a single-center retrospective study of patients who underwent device implantation and/or
with known SCN5A mutations. Genetic testing was not sent on all patients who underwent
device implantation and the testing was not uniform across patients. Therefore, we cannot
exclude the possibility of additional mutations that have not been discovered.

Conclusions and Significance

In this study, we identified an association between sinus node dysfunction, atrial arrhythmias,
elevated or fluctuating poor pacemaker capture, and SCN5A mutations (5 heterozygous and 1
homozygous) that are predicted to cause severely compromised NaV1.5 function. Our findings
underscore the importance of maintaining a high index of suspicion for genetic mutations when
working up patients with complex arrhythmias or poor pacemaker capture, especially when they
exhibit sinus node dysfunction and atrial arrhythmias. Whether these findings represent a unique
clinical manifestation of SCN5A mutations or are part of the continuum of the Brugada
phenotype and overlap syndrome warrants further investigation. Recognition of this association
can have important implications for device implantation strategies in order to improve clinical
outcomes.

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Conflict of Interest Disclosures: None

References:

AL, Jr. Congenital sick sinus syndrome caused by recessive mutations in the cardiac sodium


Table 1: The 11 patients with poor pacemaker capture out of 136 reviewed at Texas Children’s Hospital.

<table>
<thead>
<tr>
<th>Case</th>
<th>Gender</th>
<th>Age at Implant in Years and Device*</th>
<th>Phenotype</th>
<th>Sensing Issue</th>
<th>Capture Issue</th>
<th>SCN5A SNP or Mutation</th>
<th>Na_1.5†</th>
<th>PolyPhen-2</th>
</tr>
</thead>
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<tr>
<td>1</td>
<td>F</td>
<td>4.7 PM</td>
<td>SND, atrial flutter; alive</td>
<td>A</td>
<td>A+V</td>
<td>G to A substitution at the donor splice site of intron 5 (IVS5+1G&gt;A)</td>
<td>Truncation to 204 aa</td>
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<td>2</td>
<td>M</td>
<td>Newborn PM</td>
<td>SND, AT, AVB, LVNC, VT, loss of V capture; alive</td>
<td>-</td>
<td>V</td>
<td>Nucleotide substitution (5227G&gt;A)</td>
<td>G1743R (trafficking defect)</td>
<td>Probably Damaging</td>
</tr>
<tr>
<td>3</td>
<td>M</td>
<td>1.6 PM</td>
<td>SND, atrial flutter/fibrillation, VT, loss of V capture; died</td>
<td>A+V</td>
<td>A+V</td>
<td>Nucleotide substitution (4534C&gt;T)</td>
<td>R1512W (impaired fast inactivation)</td>
<td>Probably Damaging</td>
</tr>
<tr>
<td>4</td>
<td>M</td>
<td>15.6 PM</td>
<td>SND, atrial flutter; alive</td>
<td>A</td>
<td>A</td>
<td>Nucleotide substitution (3823G&gt;A)</td>
<td>D1275N (reduced I_{Na})</td>
<td>Probably Damaging</td>
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<td>5</td>
<td>F</td>
<td>11.8 ICD</td>
<td>SND, atrial flutter/fibrillation, VT; alive</td>
<td>A</td>
<td>A+V</td>
<td>Nucleotide substitution (5770G&gt;A)</td>
<td>A1924T (impaired activation)</td>
<td>Possibly Damaging or Benign</td>
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<td>12.3 ICD</td>
<td>SND, AT; alive</td>
<td>V</td>
<td>A+V</td>
<td>Nucleotide substitution (5770G&gt;A)</td>
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<td>Deletion of “C” at position 100 in exon 2 (100delC)</td>
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<td>16.0 PM</td>
<td>LVNC, LQT, AT, VT, loss of V capture; died</td>
<td>-</td>
<td>A</td>
<td>SNPs: 1673A&gt;G, 100C&gt;T</td>
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<td>Sinus pauses, intermittent AVB, A flutter; alive</td>
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*Abbreviations: PM=pacemaker; ICD=implantable cardioverter defibrillator; SND=sinus node dysfunction; A=atrial; V=ventricular; AT=atrial tachycardia; AV=atrial-ventricular; AVB=AV block; LVNC=left ventricular non-compaction; VT=ventricular tachycardia; NS=nonspecific. †Full length Na_1.5 is 2016 amino acids (aa) long. ‡Homozygous
Figure Legends:

Figure 1: Retrospective review of all patients with new pacemaker or ICD implant from 2009-2014 at Texas Children’s Hospital. A. Of these 136 patients, 11 had capture issues. Of these, 1 did not undergo genetic testing while the remaining 10 exhibited sinus node dysfunction (SND) and atrial arrhythmia (AA). B. Genetic testing of these 10 patients revealed that 7 of them (70%) harbored SCN5A mutations, 2 had multiple SCN5A single-nucleotide polymorphisms (SNPs), and 1 had a variant of unknown significant in the DES gene, which codes for desmin.

Figure 2: Conservation of the amino acids that are mutated in patients from Case 3 (A), Case 4 (B), and Case 5 (C). The left panel lists the species while the arrows point to the amino acid and position that are mutated respectively in Cases 3, 4, and 5.

Figure 3: Retrospective review of all patients with confirmed SCN5A mutations at Texas Children’s Hospital. A. The large circle represents the 27 patients with SCN5A mutations. Of these, 15 patients had either a pacemaker or implantable cardioverter defibrillator (ICD) (medium circle). Out of the 15 patients, 7 had sinus node dysfunction (SND), atrial arrhythmia (AA), and pacemaker capture issues (small circle). B. Of the 7 patients with both SCN5A mutation and SND, AA and capture issues, none had a definitive diagnosis of Brugada or long QT (LQT) syndrome. In contrast, of the patients with SCN5A mutation without capture issues, 3 had Brugada and 5 had long QT syndrome.

Figure 4: Novel SCN5A mutation and its predicted effect. This mutation is a single nucleotide
deletion at position 100 of SCN5A exon 2 (100delC), which is predicted to lead to a frame-shift of the downstream message. This introduces a premature stop codon resulting in a truncated protein product of 95 amino acids (aa) long. The wild-type product (NaV1.5) is 2016 aa long. Black boxes=exons; white boxes=introns.

**Figure 5:** Schematic of NaV1.5 encoded by SCN5A and the mutations in this study. The filled circles refer to mutations leading to NaV1.5 truncations while the open circles refer to missense mutations. The numbers inside the circles correspond to the Case numbers, except for “5” which refers to both Cases 5 and 6 since they are identical. DI-IV = domains 1-4. Figure modified from Splawski *et al.*\(^4\), with permission.
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Loss-of-Function SCN5A Mutations Associated with Sinus Node Dysfunction, Atrial Arrhythmias, and Poor Pacemaker Capture
David Y. Chiang, Jeffrey J. Kim, Santiago O. Valdes, Caridad de la Uz, Yuxin Fan, Jeffrey Orcutt, Melissa Domino, Melissa Smith, Xander H.T. Wehrens and Christina Y. Miyake

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