Reduced Na\(^+\) and higher K\(^+\) channel expression and function contribute to right ventricular origin of arrhythmias in \textit{Scn5a}+/− mice

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1. Summary

Brugada syndrome (BrS) is associated with ventricular tachycardia originating particularly in the right ventricle (RV). We explore electrophysiological features predisposing to such arrhythmic tendency and their possible RV localization in a heterozygotic \textit{Scn5a}+/− murine model. Na\(_\text{ATP}\) and protein expression were lower in \textit{Scn5a}+/− than wild-type (WT), with a further reduction in the RV compared with the left ventricle (LV). RVs showed higher expression levels of K\(_\text{ATP}\), K\(_\text{ATP}^<\), and KChIP2 in both \textit{Scn5a}+/− and WT. Action potential upstroke velocity and maximum Na\(^+\) current (\(I_{\text{Na}}\)) density were correspondingly decreased in \textit{Scn5a}+/−. These findings are predictive of a localized depolarization abnormality leading to slowed conduction. Persistent Na\(^+\) current (\(I_{\text{pNa}}\)) density was decreased in a similar pattern to \(I_{\text{Na}}\). RV transient outward current (\(I_{\text{to}}\)) density was greater than LV in both WT and \textit{Scn5a}+/−, and had larger time constants of inactivation. These findings were also consistent with the observation that AP durations were smallest in the RV of \textit{Scn5a}+/−, fulfilling predictions of an increased heterogeneity of repolarization as an additional possible electrophysiological mechanism for arrhythmogenesis in BrS.
2. Introduction

Brugada syndrome is characterized by increased clinical risks of ventricular tachycardia (VT). It has been associated with a loss of Na⁺ channel function. Clinical studies suggest that BrS is an RV disease resulting in an electrocardiographic, right precordial, ST elevation, right bundle branch block and changes specific to RV epicardial AP waveforms [1]. A range of mechanisms may contribute to this arrhythmic phenotype. In addition to potential contributions from fibrotic change and reduced connexin expression [2], electrophysiological abnormalities could result in either abnormal pNa relative to the early repolarizing I_	ext{Na}, with or without accompanying changes in I_	ext{Kr}.

The present study tests a prediction that one or more of the latter electrophysiological abnormalities might be preferentially localized to the RV at the cellular and molecular level, and thereby give rise to the arrhythmic phenotype seen at the clinical or whole heart level. It thus explores for preferential alterations in I_	ext{Na} and I_	ext{K} expression in the RV in a murine heterozygotic Scn5a⁺/− model for BrS. Previous studies have either relied on largely non-invasive experiments in humans or on a canine pharmacological wedge preparation in which the BrS phenotype was replicated by multiple pharmacological interventions. In contrast, although BrS is a genetically heterogeneous condition, up to 30 per cent of patients have mutations in the Scn5a gene [6]. Thus far, it is the only gene that has been extensively studied in its respective threshold cycle (CT) using a standard curve. All experiments were performed in duplicate. The samples were then centrifuged at 2000 × g for 10 min with a pestle and mortar. They were then homogenized using a Dounce homogenizer (Jencons, Lutterworth, UK) in ice-cold Krebs–Henseleit buffer containing: 119 mM NaCl, 25 mM NaHCO₃, 4 mM KCl, 1.2 mM KH₂PO₄, 1 mM MgCl₂, 1.8 mM CaCl₂, 10 mM glucose and 2 mM sodium pyruvate (pH 7.4). The ventricles were removed and the LV and RV free walls were isolated and flash-frozen. The tissues were ground in liquid nitrogen with a pestle and mortar. They were then homogenized using a Dounce homogenizer (Jencons, Lutterworth, UK) in ice-cold lysis buffer containing: 50 mM Tris pH 8.0, 150 mM NaCl, 1 per cent NP40, 0.2 per cent SDS, 0.5 per cent Na deoxycholate, 1 mM EDTA and complete protease inhibitor cocktail (Roche, Burgess Hill, UK), and rotated for 1 h at 4°C. The samples were then centrifuged at 2000g for 10 min

3. Methods

3.1. Experimental animals

Mice aged three to five months were obtained from breeding pairs of heterozygote Scn5a⁺/− and WT inbred 129/sv mice. Hearts from all groups were subject to identical experimental procedures, permitting their comparison. WT mice were initially supplied by Harlan (UK), and Scn5a⁺/− mice were originally generated from these as described previously [12]. Mice were killed by cervical dislocation, in compliance with the UK Animals (Scientific Procedures) Act 1986. The investigation conforms to the Guide for the Care and Use of Laboratory Animals and was performed under licences approved by local ethical committees and issued by the UK Home Office.

3.2. Real time reverse transcriptase-PCR

To quantify changes in the mRNA expression levels of Na⁺ channel α-subunits and a range of K⁺ channels in murine hearts, real time reverse transcriptase-PCR (RT-PCR) experiments were performed on an ABI 7500 Fast Cycler (Applied Biosystems, Warrington, UK). Total RNA was isolated from the LV and RV of WT and Scn5a⁺/− mice (n = 4 each) using a Qiagen RNAeasy kit. Excised tissues were stored in RNAlater (Ambion, Warrington, UK) to maintain the integrity of the RNA before isolation. The total RNA was reverse transcribed into cDNA using random hexamer primers and a SuperScript III kit (Invitrogen, Paisley, UK). Oligos for Scn5a were fluorescent amidite (FAM) labelled (Applied Biosystems). All experiments were performed in duplicate.

The number of copies of mRNA was calculated from its respective threshold cycle (Ct) using a standard curve. The 2⁻∆∆CT method [13] was used to calculate gene expression relative to the expression of the housekeeper gene, glyceraldehyde-3-phosphate dehydrogenase (GAPDH). These values were then normalized to the mean Na,1.5 gene expression in the LV of the WT mice.

3.3. Western blots

Hearts were excised and placed in ice-cold Krebs–Henseleit buffer containing: 119 mM NaCl, 25 mM NaHCO₃, 4 mM KCl, 1.2 mM KH₂PO₄, 1 mM MgCl₂, 1.8 mM CaCl₂, 10 mM glucose and 2 mM sodium pyruvate (pH 7.4). The ventricles were removed and the LV and RV free walls were isolated and flash-frozen. The tissues were ground in liquid nitrogen with a pestle and mortar. They were then homogenized using a Dounce homogenizer (Jencons, Lutterworth, UK) in ice-cold lysis buffer containing: 50 mM Tris pH 8.0, 150 mM NaCl, 1 per cent NP40, 0.2 per cent SDS, 0.5 per cent Na deoxycholate, 1 mM EDTA and complete protease inhibitor cocktail (Roche, Burgess Hill, UK), and rotated for 1 h at 4°C. The samples were then centrifuged at 2000g for 10 min
and the supernatant kept. The pellet was resuspended in lysis buffer and recentrifuged at 2000g for a further 10 min, and the second supernatant added to the first. The combined supernatants were then ultracentrifuged at 100,000g for 1 h, and the pellet suspended in buffer containing: 4 mM HEPES, 320 mM sucrose and complete protease inhibitor cocktail. The protein concentration of each sample was established using a Pierce bicinchoninic acid assay kit (Thermo Scientific, Rockford, USA), using microplate assays in triplicate.

All equipment and consumables mentioned in this paragraph are from Invitrogen unless otherwise stated. Protein samples were mixed with 4x NuPAGE LDS sample buffer, 10x β-mercaptoethanol and H2O2, and were heated at 70°C for 10 min. XCell SureLock Mini-Cells were used to run gel electrophoresis. One of two sets of electrophoresis conditions were used depending on whether the protein to be analysed was of high or low molecular weight. For high molecular weight proteins, electrophoresis was carried out using NuPAGE Novex 3 to 8 per cent Tris-Acetate gels at 30 V for 90 min, and then at 140 V overnight at 4°C with NuPAGE Tris-Acetate running buffer and NuPAGE antioxidant, alongside a Hi-Mark pre-stained high-molecular-weight protein standard. For low-molecular-weight proteins, electrophoresis was carried out using NuPAGE Novex 4 to 12 per cent Bis-Tris gels at 140 V for 2 h with NuPAGE MES SDS running buffer and NuPAGE antioxidant, alongside a pre-stained protein standard. Twenty mM of total protein was loaded in each lane. Protein bands were transferred onto polyvinylidene difluoride (PVDF) membranes by an XCell II Blot Module, with NuPAGE transfer buffer, 10 per cent methanol and NuPAGE antioxidant, at 100 mA overnight at 4°C. Membranes were blocked for 5 h in 5 per cent BSA-phosphate-buffered saline (PBS)—1 per cent Tween, prior to overnight incubation at 4°C with primary antibody. Antibodies used were to Na+,K+, ATP levels were calculated using IMAGEJ (NIH, Bethesda, USA), and plots drawn of protein expression normalized to either heavy chain myosin or GAPDH expression.

3.4. Ventricular myocyte isolation

Myocytes were isolated as previously described [14]. Mouse hearts were rapidly removed and placed in ice-cold normal Tyrode (NT) solution containing: 140 mM NaCl, 6 mM KCl, 1 mM MgCl2, 1 mM CaCl2, 10 mM glucose, 10 mM HEPES, pH adjusted to 7.4 with 2 M NaOH. The ascending aorta was cannulated with a blunted 21 g needle. The heart was then retrogradely perfused for 2–3 min at 37°C with NT solution followed by a further 5 min in low Ca2+ (LC) solution containing: 120 mM NaCl, 5.4 mM KCl, 5 mM MgSO4, 0.045 mM CaCl2, 5 mM sodium pyruvate, 20 mM glucose, 20 mM taurine, 10 mM HEPES, 5 mM nitroliotric acid, bubbled with 100% O2: pH 6.96. The solution was changed to an enzyme solution (ES) containing: 120 mM NaCl, 5.4 mM KCl, 5 mM MgSO4, 0.2 mM CaCl2, 5 mM sodium pyruvate, 20 mM glucose, 20 mM taurine, 10 mM HEPES, bubbled with 100% O2: pH 7.4 with addition of protease (4 U ml⁻¹, Sigma) for 1 min, followed by perfusion for 8 min with ES solution containing collagenase (0.8 mg ml⁻¹; Worthington) and hyaluronidase (0.5 mg ml⁻¹, Sigma). At this stage, the heart was swollen and flaccid but with anatomy still intact. The free walls of the LV and RV were then separated and cut into small pieces using a scalpel blade. The tissue was then agitated for a further 5 min in ES solution. This suspension was gently triturated using a pipette, filtered and resuspended in ES solution. Myocytes were stored in ES solution at room temperature and used within 7–8 h of isolation. Isolated myocytes were checked under a light microscope to confirm cell integrity. Live myocytes appeared as smooth and rod-shaped cells.

3.5. Electrophysiology measurements

3.5.1. Measurement of action potentials

Cells were superfused with NT solution at 37°C and studied using a perforated patch technique using an Axon 2B amplifier (Axon Instruments, CA, USA). The pipette resistance was 1–2 MΩ, and the pipette-filling solution contained: 125 mM KCl, 20 mM KCl, 10 mM NaCl, 10 mM HEPES, 5 mM MgCl2, pH 7.2 with KOH, plus amphotericin-B (final concentration 240 μg ml⁻¹). APs were measured in current-clamp mode after stimulation at 1, 3 and 5 Hz using a 1 ms, 1.2–1.4 nA pulse. The APs were measured and averaged over 10 consecutive beats. Traces were then averaged with reference to the stimulation signal, and times to 50 per cent (APD50), 70 per cent (APD70) and 90 per cent repolarization (APD90) were taken as a measure of the AP duration for comparison between groups. Maximum upstroke velocity was measured as the maximum gradient of the upstroke of the AP waveform.

3.5.2. Current density measurements

For all current density measurements, the whole-cell configuration of the patch-clamping technique was employed using a Multiclamp 700A amplifier (MultiClamp 700A, Molecular Devices, USA), and averaged over 30 consecutive beats. Band intensity was calculated using ImageJ (NIH, Bethesda, USA), and was assessed as [14]. Mouse hearts were rapidly removed and placed in ice-cold normal Tyrode (NT) solution containing: 140 mM NaCl, 6 mM KCl, 1 mM MgCl2, 1 mM CaCl2, 10 mM glucose, 10 mM HEPES, pH adjusted to 7.4 with 2 M NaOH. The ascending aorta was cannulated with a blunted 21 g needle. The heart was then retrogradely perfused for 2–3 min at 37°C with NT solution followed by a further 5 min in low Ca2+ (LC) solution containing: 120 mM NaCl, 5.4 mM KCl, 5 mM MgSO4, 0.045 mM CaCl2, 5 mM sodium pyruvate, 20 mM glucose, 20 mM taurine, 10 mM HEPES, 5 mM nitroliotric acid, bubbled with 100% O2: pH 6.96. The solution was changed to an enzyme solution (ES) containing: 120 mM NaCl, 5.4 mM KCl, 5 mM MgSO4, 0.2 mM CaCl2, 5 mM sodium pyruvate, 20 mM glucose, 20 mM taurine, 10 mM HEPES, bubbled with 100% O2: pH 7.4 with addition of protease (4 U ml⁻¹, Sigma) for 1 min, followed by perfusion for 8 min with ES solution containing collagenase (0.8 mg ml⁻¹; Worthington) and hyaluronidase (0.5 mg ml⁻¹, Sigma). At this stage, the heart was swollen and flaccid but with anatomy still intact. The free walls of the LV and RV were then separated and cut into small pieces using a scalpel blade. The tissue was then agitated for a further 5 min in ES solution. This suspension was gently triturated using a pipette, filtered and resuspended in ES solution. Myocytes were stored in ES solution at room temperature and used within 7–8 h of isolation. Isolated myocytes were checked under a light microscope to confirm cell integrity. Live myocytes appeared as smooth and rod-shaped cells.

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3.5.2. Current density measurements

For all current density measurements, the whole-cell configuration of the patch-clamping technique was employed using a Multiclamp 700A amplifier (MultiClamp 700A, Molecular Devices, USA), Borosilicate glass pipettes (0.86 mm outer diameter; Harvard Apparatus, Kent, UK) were used, with a resistance of 1.5–2.5 MΩ. All the experiments were performed at 37°C, apart from those measuring the fast Na⁺ current, which were performed at a constant room temperature of 22°C. The rate of superfusion was 2–3 ml min⁻¹. Miniature solenoid valves (The Lee Company, Essex, UK) were used to produce fast changes in superfusate. All the recordings were done after 3 min of membrane rupture, after which time pilot experiments demonstrated stable voltage dependency of currents. Each set of measurements took approximately 5 min to complete. Capacitance and 80 to 90 per cent series resistance
compensation was performed prior to each recording. Only cells with initial resting potentials between −70 and −80 mV, initial seal resistances greater than 5 GΩ with stable residual series resistances less than 1 MΩ, and low leakage currents (less than 0.1 nA at −120 mV) were studied. Cell stability in the course of the experiments was further followed by comparing transients from the same (largest) voltage steps at successive times during the protocol.

The currents obtained were filtered at a 10 kHz high-frequency cut-off (four pole Bessel filter) and digitized at 100 kHz using pCLAMP software.

The membrane capacitance (in picofarads) of each cell was initially measured by integrating the area under a capacitative transient induced by a 10 mV hyperpolarizing clamp step (from −80 to −90 mV) and dividing this area by the voltage step. There were no significant differences in mean cell capacitance between groups (122.4 ± 6 pF in WT LV, 119.5 ± 7 pF in WT RV, 129.3 ± 6 pF in Scn5a+/− LV and 126.6 ± 9 pF in Scn5a+/− RV; n = 36; p > 0.05). All the currents were then normalized to the whole-cell capacitance for cross-group comparisons.

3.6. Na⁺ current, Iₙa

The fast Na⁺ current Iₙa was measured using a technique previously described [15]. Pipettes were filled with: 70.26 mM CsCl, 24.74 mM Cs-aspartate, 10 mM HEPES, 1 mM Na₂ATP, 2 mM MgATP, 1.37 mM MgCl₂ and 10 mM Cs-BAPTA, adjusted to pH 7.2 using CsOH. Cells were initially perfused with NT solution containing 200 µM Cd²⁺. To achieve optimum voltage control, Na⁺ currents were recorded in a low [Na⁺] solution. This solution contained: 10 mM NaCl, 140 mM CsCl, 10 mM glucose, 10 mM HEPES, 1 mM MgCl₂ and 1 mM CaCl₂, adjusted to pH 7.4 using CsOH. Na⁺ currents were initially measured before and after perfusion with 10 mM tetrodotoxin (TTX), and the TTX sensitive current calculated; however, pilot studies showed that under our experimental conditions, TTX eliminated the Na⁺ current calculated; however, pilot studies demonstrated evidence of inadequate voltage control (e.g. a ‘threshold phenomenon’ near the voltage range for Na⁺ channel activation) data were discarded. The absence of crossover of the current traces as the current magnitude was changed and the shape of the current–voltage (I−V) relationship remaining unchanged for difference peak current values further suggest adequate voltage control [16,17].

First, the voltage dependence of activation was studied by applying depolarizing pulses of 250 ms duration over a series of voltage steps from a fixed holding voltage of −120 mV, in 5 mV increments, to +20 mV. The peak current densities (pA/pF) were calculated by dividing the peak currents amplitude by Cm and the I−V relationship plotted. The values of peak Na⁺ conductance (gₙa) were determined from the equation

\[ g_{Na} = \frac{I_{Na}}{V_m - E_{Na}} \]

as first described by Hodgkin et al. [18] as a linear approximation of the Goldman–Hodgkin–Katz equation and subsequently used extensively in patch clamp analysis of cardiac myocytes [15,19–21], where Iₙa is the peak current density and the Eₙa is the [Na⁺] reversal potential. The gₙa values calculated from each myocyte were normalized against the corresponding peak gₙa value. When plotted against the test-pulse voltage, the data assumed a sigmoid function, increasing with membrane potential.

The voltage dependence of steady-state inactivation was also examined using conditioning pre-pulses of 500 ms duration. These variable pre-pulses were applied over a voltage range from −140 to −30 in 5 mV increments. Each pre-pulse was followed by a constant test pulse to −20 mV at a duration of 20 ms. Normalized currents were then plotted against the pre-pulse voltage. The data assumed a sigmoid function, decreasing with membrane potential. Both inactivation and activation data were fitted with a single Boltzmann function, as used extensively in previous studies [15,20–24], of the form:

\[ y = \left[ 1 + \exp \left( \frac{s(V_m - V_{1/2})}{k} \right) \right]^{-1} \]

For activation curves, the y-values correspond to the normalized conductance values and constant s equals −1. For inactivation curves, the y-values correspond to the normalized current values and constant s equals +1. This yields the half-maximal voltage (V₁/₂) and exponential slope factor (k) values.

3.6.1. Persistent Na⁺ current

The persistent Na⁺ current (Iₚₙa) was measured using a technique previously described [25]. Cells were superfused at 37°C with modified NT solution, with K⁺ substituted with Cs⁺, and containing 10 µM nifedipine (Sigma, Poole, UK), 100 µM strophantidin (Sigma) and 30 µM niflamic acid (Sigma). The pipette filling solution contained: 115 mM Cs-aspartate, 20 mM TEA-Cl, 10 mM EGTA, 10 mM HEPES, 5 mM MgATP, pH 7.2. From a holding potential of −100 mV, increasing voltage (20 mV) steps from −80 mV to +20 mV for 1 s were applied with a cycle length of 5 s. The protocol
was repeated in the presence of 20 μM TTX (Alomone Labs., Israel), and the TTX-sensitive current was analysed. \( I_{pNa} \) density was taken as the average current between 30 and 80 ms after depolarization and normalized to cell capacitance. This period was chosen because this is approximately the range of APDs measured in our experiments.

### 3.6.2. Transient outward current, \( I_{to} \)

The transient outward current \( I_{to} \) was measured using a technique previously described [26]. Cells were superfused with NT solution containing 200 μM Cd\(^{2+}\) and the pipette-filling solution contained: 120 mM K-glutamate, 10 mM KCl, 2 mM MgCl\(_2\), 10 mM HEPES, 5 mM EGTA, 2 mM Mg-ATP, pH 7.2 with KOH at 37°C. Pipette resistance was 1.5–2.5 M\(\Omega\).

\( I_{to} \) was elicited by increasing voltage steps (1 s duration) from -40 mV to +70 mV with a cycle length of 5 s in increments of 10 mV. In these studies, the transient outward current was separated from the other currents by the following procedures. The holding potential was held at -40 mV to inactivate the sodium current. \( I_{Ca} \) was largely blocked by 0.2 mM CdCl\(_2\) included in the recording solution. EGTA was included in the pipette solution to minimize Ca\(^{2+}\)-induced inactivation. \( I_{to} \) was taken as the peak of the transient outward current \( I_{to,peak} \) from which the steady-state current \( I_{ss} \) had been subtracted. While \( I_{ss} \) may also demonstrate inactivation, studies have shown that this plays a relatively minor role in repolarization in the murine ventricle [27,28]. As for \( I_{Na} \), the \( I\text{-}V \) relationship was plotted, as well as the voltage dependence of activation as determined from conductance calculations, and fitted with a single Boltzmann distribution.

The two time constants were plotted as a function of voltage. Steady-state inactivation parameters of \( I_{to,peak} \) were obtained with a double-pulse protocol using 1 s prepulses to voltages from -100 to +20 mV from a holding potential of -100 mV. Peak currents at the test pulse to +40 mV were normalized to the maximal transient outward current elicited by the test pulse and fitted by a single Boltzmann distribution.

### 3.7. Statistical procedures

Patch clamp measurements were made from a set of 8 WT and 8 Scn5a\(^{+/\text{--}}\) mice, with \( n \) numbers representing the total number of cells studied. 4 WT and 4 Scn5a\(^{+/\text{--}}\) mice were each used for RT-PCR experiments and a further four each for Western blot experiments. All electrophysiological parameters and expression levels of protein and mRNA were expressed as mean ± s.e.m. values. Their significant differences were analysed using ANOVA applied to all groups with post-hoc Tukey’s honestly significant difference tests then applied to paired groups. A \( p \)-value of less than 0.05 was the criterion for significance.

### 4. Results

#### 4.1. Real-time reverse transcriptase-PCR

Na\(_{1.5}\) mRNA expression was significantly lower in Scn5a\(^{+/\text{--}}\) hearts than in WT, as previously reported [22] (figure 1). While this was true in both the LV and RV, there was a bigger drop in the RV. While in the LV expression fell to approximately two-thirds of that in the WT, in the RV it fell to about one-third of that in the WT. Interestingly, in the WT there was significantly higher expression of Na\(_{1.5}\) mRNA in the RV than LV, while in the Scn5a\(^{+/\text{--}}\) hearts, there was no significant difference between mRNA expression in the two ventricles.
Regarding mRNA expression for K⁺ channels in the heart, we found no significant differences either between LV and RV, or between WT and Scn5a+/− hearts for Kir2.1, Kv1.4 and Kv1.5. However, there were significantly higher mRNA expression levels of Kv4.2 in RVs in both WT and Scn5a+/− hearts, although there was no difference between WT and Scn5a+/−. Kv4.3 showed a small increase in WT RV compared with the WT LV or Scn5a+/− RV, but there were no differences between LV WT or Scn5a+/−, nor between LV and RV Scn5a+/−. Lastly, there was higher mRNA expression of KChIP2 in WT than Scn5a+/− RV, and in the RV than LV, of both WT and Scn5a+/− hearts.

4.2. Western blots

Western blots of Na,1.5 demonstrated a single band at around 220 kDa (predicted weight 227 kDa; figure 2a). There was no significant difference in the Na,1.5 expression found between the LV and RV of WT hearts (figure 2b). As expected, Scn5a+/− hearts had significantly reduced Na,1.5 expression, but there was a further difference between LV and RV, with approximately 30 per cent reduction in the LV and approximately 50 per cent reduction in the RV when compared with WT, resulting in a significantly lower expression of Na,1.5 in the RV compared with the LV of Scn5a+/− hearts. For K⁺ channels, there were no significant differences in protein expression either between LV and RV or WT and Scn5a+/− hearts for Kir2.1, Kv1.4 or Kv1.5 (figure 2c,d). However, there were significantly higher protein levels in the RV than LV for both WT and Scn5a+/− hearts for Kv4.2, Kv4.3 and KChIP2, with no difference between WT and Scn5a+/− hearts.

4.3. Electrophysiology

APDs and I_{Na}, I_{PNa} and I_{v} densities were measured using patch clamp techniques. APs from all groups demonstrated a triangular AP morphology, with a resting membrane potential of around −70 mV and with an overshoot to around +40 mV, in agreement with previous studies in WT mice [29].
4.4. Scn5a+/−: right ventricle action potentials have shorter durations and lower maximum upstroke velocities compared with respective left ventricles and with wild-type

The maximum upstroke velocity (dV/dt_{max}) of the APs, measured as the maximum gradient of the AP upstroke, can be used to give an indication of Na\(^+\) channel function [30]. dV/dt_{max} was reduced in Scn5a+/− hearts compared with WT in both LV and RV myocytes at all pacing frequencies (figure 3). The dV/dt_{max} values were not significantly different between LV and RV in WT myocytes, but were significantly reduced in RV compared with LV in Scn5a+/− myocytes. There were no significant differences overall in the amplitude of the APs between any group.

At all three pacing frequencies, APD_{50} was significantly lower in the RV compared with the LV of Scn5a+/− myocytes, but not in WT. The APD_{50} of RV Scn5a+/− myocytes was also significantly less than that of RV WT myocytes, while there were no differences between the respective LVs. Similar results were obtained for the APD_{90} measurements. These myocyte data correlate well with previous studies from AP recordings using monophasic AP electrodes in Langendorff perfused hearts [31].

4.5. Scn5a+/−: right ventricle myocytes have lower I_{Na} than the respective left ventricle and in wild-type

Na\(^+\) currents were initially measured before and after perfusion with 10 \(\mu\)M TTX, and the TTX-sensitive current calculated; however, pilot studies showed that under our experimental conditions, TTX eliminated the entire current being studied, and therefore showed that we were specifically recording the Na\(^+\) current without the need for TTX subtraction (figure 4). There were no significant differences in mean capacitance of myocytes between any of the groups.

Values of maximum Na\(^+\) current density (I_{Na, max}) from WT myocytes were in agreement with those from previous studies [23]. There were no significant differences in I_{Na, max} or I–V relationship between LV and RV myocytes from WT hearts (figure 5a–c). I_{Na, max} was significantly less in both LV and RV of Scn5a+/− hearts than in WT; this was also reflected in differences in the I–V relationship. In addition, in Scn5a+/− hearts Na\(^+\) current density was reduced in the RV compared with the LV. Overall, I_{Na, max} in the RV of Scn5a+/− hearts was around 50 per cent that of RV WT, while in the Scn5a+/− LV it was around 70 per cent of WT LV.

Activation curves yielded parameters of voltage dependence of activation similar to previous studies [12,22,23]. There were no significant differences in these parameters between any of the groups studied (figure 5d and table 1). The inactivation curve was shifted to more negative values in the Scn5a+/− hearts compared with WT for both RV and LV, and for the RV was further shifted to more negative values. In addition, in Scn5a+/− hearts Na\(^+\) current density was reduced in the RV compared with the LV. Overall, I_{Na, max} in the RV of Scn5a+/− hearts was around 50 per cent that of RV WT, while in the Scn5a+/− LV it was around 70 per cent of WT LV.

4.6. Lower I_{Na} in Scn5a+/−: right ventricle myocytes compared with respective left ventricle and with wild-type

We proceeded to measure the persistent Na\(^+\) current, I_{pNa}. Following a similar pattern to I_{Na}, maximum I_{pNa} was...
Figure 4. (a) Example trace of an $I_{Na}$ current from a myocyte from the LV of a WT heart, before addition of 10 μM TTX, and then with TTX subtraction. (b) $I-V$ curve for the two conditions (squares, control; circles, TTX subtraction), with no significant differences ($n = 4$).

Figure 5. $I_{Na}$ measurements. (a) Example traces of Na$^{+}$ currents from myocytes from the LV and RV of WT and Scn5a+/− hearts. The traces have been normalized to cell capacitance, and the protocol is shown in the inset. (b) $I-V$ relationship of Na$^{+}$ current in myocytes from the four groups ($n = 17$). (c) Maximum $I_{Na}$ density of each group ($n = 17$). (d) Activation and (e) inactivation curves in myocytes from the four groups, with Boltzmann fits ($n = 17$ for activation, $n = 12$ for inactivation). Significant differences: asterisks (*), effect of genotype; hashes (#) effect of cardiac ventricle. Note in (b), upper asterisk denotes differences in genotype for RV, and lower asterisk for LV.
Figure 6. \(I_{\text{Na}}\) measurements. (a) Example traces from a test pulse to \(-20\) mV from the LV and RV of WT and \(\text{Scn5a}^{+/+}\) myocytes. The vertical lines denote 30 and 80 ms after depolarization. (b) \(I–V\) relationship of mean \(I_{\text{Na}}\) from 30 to 80 ms after depolarization in myocytes from each group. (c) Bar graph of maximum \(I_{\text{Na}}\) density averaged from 30 to 80 ms after depolarization. Significant differences: asterisks (*), effect of genotype; hashes (#), effect of cardiac ventricle (n = 10 for each group).

Table 1. Parameters of activation and inactivation from Boltzmann fits (millivolts) to the curves produced from \(I_{\text{Na}}\) in myocytes from the LV and RV of WT and \(\text{Scn5a}^{+/+}\) hearts (n = 17 for activation; n = 12 for inactivation for each group).

|       | \(V_{1/2}\)     | \(k\)       | \(V_{1/2}\)     | \(k\)       |
|-------|----------------|------------|----------------|------------|
| WT LV | \(-58.88 \pm 2.85\) | \(3.65 \pm 0.66\) | \(-74.64 \pm 1.83\) | \(7.95 \pm 0.62\) |
| WT RV | \(-60.42 \pm 3.04\) | \(4.14 \pm 0.42\) | \(-75.35 \pm 1.72\) | \(7.21 \pm 0.60\) |
| \(\text{Scn5a}^{+/+}\) LV | \(-64.61 \pm 2.96\) | \(3.44 \pm 0.78\) | \(-84.16 \pm 1.53^*\) | \(6.56 \pm 0.71\) |
| \(\text{Scn5a}^{+/+}\) RV | \(-63.69 \pm 3.40\) | \(5.46 \pm 0.93\) | \(-91.12 \pm 1.64^{a,b}\) | \(6.61 \pm 0.74\) |

\(^a\)Significant effect of genotype.
\(^b\)Significant effect of cardiac ventricle.

4.7. Higher \(I_{\text{Na}}\) in right ventricle than left ventricle of wild-type and \(\text{Scn5a}^{+/+}\) myocytes

Lastly, we measured inactivating K\(^+\) current by subtracting the sustained current \(I_{\text{Na}}\) from the point of depolarization onwards in raw traces (figure 7a). The inactivating K\(^+\) current was taken to represent \(I_{\text{Na}}\) max, as, while more slowly inactivating K\(^+\) currents do exist in the mouse ventricle, they are relatively small [27]. There were no significant differences in \(I_{\text{Na}}\) amplitude between any of the groups. \(I_{\text{Na}}\) max was significantly greater in the RV than LV of both \(\text{Scn5a}^{+/+}\) and WT, which was also reflected in differences in the \(I–V\) relationship (figure 7b,c). There were no significant differences in \(I_{\text{Na}}\) max or the \(I–V\) relationship between WT and \(\text{Scn5a}^{+/+}\) myocytes from either LV or RV.

Analysis of the kinetics of inactivation through fitting to a double exponential decay function demonstrated three components of the K\(^+\) current, which are likely to represent \(I_{\text{Na}}, I_{\text{Ka}}\) and \(I_{\text{Slow}}\) respectively [32]. The amplitude of \(I_{\text{Fast}}\) was approximately 10 times that of \(I_{\text{Slow}}\), supporting evidence that \(I_{\text{Na}}\) is the dominant inactivating K\(^+\) current in the mouse ventricle [27]. Figure 7d shows the time constants of decay of the two inactivating components from myocytes from all groups. There was no clear voltage dependence, similar to findings in other studies [32].

The RV showed values of \(\tau_{\text{Fast}}\) and \(\tau_{\text{Slow}}\) that were higher than their respective values in the LV in both the WT and \(\text{Scn5a}^{+/+}\). These time constants were then higher in the \(\text{Scn5a}^{+/+}\) than WT in the RV for all voltages for \(\tau_{\text{Fast}}\) and at more positive voltages for \(\tau_{\text{Slow}}\). In contrast, the LV of both WT and \(\text{Scn5a}^{+/+}\) showed similar values of \(\tau_{\text{Fast}}\) and \(\tau_{\text{Slow}}\).

There were no significant differences in voltage dependence of activation (figure 7e and table 2). Voltage dependence of inactivation was shifted to more negative values in the RV compared with LV of both WT and \(\text{Scn5a}^{+/+}\) hearts, but there was a shift to more positive values for \(\text{Scn5a}^{+/+}\) myocytes compared with WT (figure 7f and table 2).

5. Discussion

These experiments use a \(\text{Scn5a}^{+/–}\) murine model to test a hypothesis that localizes potentially arrhythmogenic
electrophysiological mechanisms to the RV. This system has previously been used to model BrS, and mapping studies have demonstrated slowed conduction and repolarization heterogeneities localized to the RV leading to re-entrant polymorphic VT originating specifically in the RV [3,8]. Our findings add to (i) reports associating the $\text{Scn5a}^{+/2}$ mutation with reduced Na$^+$ current expression [12], and (ii) previous studies in WT mice that demonstrated higher $\text{Scn5a}$ mRNA and protein expression and Na$^+$ channel availability in endocardium than epicardium [33]. Neither study made comparisons between RV and LV.

Table 2. Parameters of activation and inactivation from Boltzmann fits (millivolts), and the time constants of inactivation from the double exponential fit to the curves produced from $I_{to}$ in myocytes from the LV and RV of WT and $\text{Scn5a}^{+/2}$ hearts ($n = 14$ for each group).

|          | activation | inactivation |
|----------|------------|--------------|
|          | $V_{1/2}$  | $k$          |
|          | $V_{1/2}$  | $k$          |
| WT LV    | $-11.06 \pm 2.54$ | $14.26 \pm 3.24$ | $-42.72 \pm 1.86$ | $9.18 \pm 1.10$ |
| WT RV    | $-9.07 \pm 2.85$ | $17.24 \pm 3.21$ | $46.40 \pm 1.72^{a}$ | $7.80 \pm 0.92$ |
| $\text{Scn5a}^{+/2}$ LV | $-9.58 \pm 3.20$ | $14.15 \pm 3.95$ | $-37.13 \pm 0.91^{b}$ | $11.30 \pm 0.70$ |
| $\text{Scn5a}^{+/2}$ RV | $-15.57 \pm 3.44$ | $20.69 \pm 3.55$ | $-42.98 \pm 1.01^{a,b}$ | $10.15 \pm 0.73$ |

*aSignificant effect of cardiac ventricle.

*bSignificant effect of genotype.
Our comparisons have fulfilled the predictions of an electrophysiological hypothesis involving a reduced \( I_{Na} \) mechanism. Thus, while the patch-clamp experiments demonstrated that WT myocytes possessed similar peak \( I_{Na} \) magnitudes in the RV and LV, these were decreased in both the RV and LV in the \( Scn5a+/- \) but with a greater reduction in RV than LV. Yet Na\(^+\) conductance activation curves were indistinguishable. These findings were also compatible with two underlying mechanisms acting upon Na\(^+\) channel function. First, there could be differing Na\(^+\) channel inactivation properties in the RV and LV between the WT and the \( Scn5a+/- \). Thus, although WT RV and LV showed similar inactivation curves, both the RV and LV of the \( Scn5a+/- \) showed negative shifts in inactivation properties, with the RV showing the greatest alterations. Second, the electrophysiological results implicated reductions in expression of otherwise normal Na\(^+\) channels. Thus, the Na\(^+\) current reductions took place even at holding voltages at which there was no significant inactivation. Furthermore, the current reductions exactly paralleled the corresponding patterns of protein expression, corroborating the latter mechanism. So did patterns of mRNA expression, apart from suggesting a relative translational upregulation in the LV relative to the RV \( Scn5a+/- \). These findings precisely correlated with the outcomes of similar AP (dV/dt)\(_{max}\) values in the RV and LV in the WT, which were reduced in \( Scn5a+/- \) to a greater extent in the RV than the LV.

Our studies also showed a reduction in \( I_{pNa} \) in the RV of \( Scn5a+/- \) hearts, which may well contribute to the shorter APDs in this region. APD values were correspondingly shorter in \( Scn5a+/- \) RV than WT RV and than \( Scn5a+/- \) LV. Other groups have reported a gradient of expression of \( I_{pNa} \) across the ventricular wall [5]. Although the physiological significance in humans is still unclear, the \( I_{pNa} \) gradient is thought to be important in the pro- and anti-arrhythmic effect of class 1 agents. This is the first study to show heterogeneity of \( I_{pNa} \) between ventricles in any model, and to contribute to increased repolarization heterogeneity.

Differential K\(^+\) channel expression has also been implicated in the arrhythmogenic mechanism in BrS. In murine hearts, K\(^+\) currents include the readily separable non-inactivating \( I_{Ks} \) component. They also include inactivating components—mainly the fast inactivating \( I_{Ko} \) and slow inactivating \( I_{Ko,s} \). These could be separated through their fast (t\(_{fast}\)) and slow (t\(_{slow}\)) time constants. As described on previous occasions [27,28], \( I_{Ko} \) appeared to play a relatively minor role in repolarization in the murine ventricle, giving a slow current \( I_{Ko,s} \) whose magnitude was less than 10 per cent that of the remaining \( I_{Kfast} \). Thus, by far the greater part of the outward current observed in the present experiments appeared to arise from \( I_{Ko} \).

\( I_{Ko} \) channels control a prominent repolarizing current, the Ca\(^2+\)-independent transient outward K\(^+\) current [34]. While in humans and large mammals, both fast (\( I_{Ko,f} \)) and slow (\( I_{Ko,s} \)) channels are expressed throughout the heart, in the mouse heart \( I_{Ko} \) is present only in the interventricular septum, and thus the focus of this study is on \( I_{Ko}^{11} \), for which the molecular correlates are K\(_{4.3}/K_{4.2} \) [35]. \( I_{Ko} \) has thus been well documented to be expressed in a heterogeneous manner throughout the ventricles of mammalian hearts, including mice, in concordance with known AP durations [32,36,37]. The present experiments demonstrated larger peak \( I_{Ko} \) magnitudes in the RV than the LV. However, there was no particular difference in this peak between \( Scn5a+/- \) and the WT. Furthermore, there were no differences in the voltage dependence of activation of \( I_{Ko} \) in RV or LV whether \( Scn5a+/- \) or WT. The voltage-dependence of \( I_{Ko} \) was slightly shifted to more negative values in the RV than the LV in both WT and \( Scn5a+/- \), which would potentially reduce its repolarizing effect in parallel with a reduced RV expression of \( I_{Ko} \). However, it was shifted to more positive values in \( Scn5a+/- \) than WT, which would accentuate the imbalance between repolarizing and depolarizing currents. Furthermore, the RV showed slower inactivation kinetics than did the LV in both WT and \( Scn5a+/- \), with a further slowing in the \( Scn5a+/- \).

In line with previous studies in mixed ventricular lysates [22], \( K_{m} \) demonstrated a similar mRNA and protein expression level greater in the RV than the LV, but with no difference between WT and \( Scn5a+/- \) in direct correlation with their peak current magnitudes. \( K_{4.3} \) showed a less clear-cut mRNA expression pattern, with small increases in WT RV compared with the remaining groups (in agreement with earlier results), but showed a similar protein expression pattern to \( K_{4.2} \). The disjunction between current magnitudes and mRNA expression would be consistent with differential KChIP2 expression, in turn controlling expression of the \( I_{Ko} \) channel. Our study supports this hypothesis in showing higher mRNA and protein expression of KChIP2 in the RV than LV of WT and \( Scn5a+/- \) hearts, which could act to increase RV expression of \( K_{4.3} \).

These findings thus directly explain changes in maximum \( I_{Na} \) and \( I_{Ko} \) In terms of altered protein expression. The mechanisms of the altered voltage dependence of the Na\(^+\) and K\(^+\) channel inactivation and K\(^+\) channel current decay particularly in the \( Scn5a+/- \) RV are unknown, but could possibly be due to changes in binding to regulatory proteins. For example, regulation of \( Na_{1.5} \) is complex, involving interactions with β-subunits and multiple signalling proteins including cytoskeletal molecules such as ankylins and synthrophins, regulatory kinases and phosphatases, components of cellular trafficking and extracellular matrix molecules, as part of large, multi-protein complexes [38]. Thus, in mice, over-expressing CaMKII, which demonstrates slowed conduction and VT, CaMKII-dependent phosphorylation of \( Na_{1.5} \) shifts the steady-state inactivation curve towards negative voltages in a Ca\(^2+\)-dependent manner [39]. Experiments on neonatal rat ventricles and heterologous cells have demonstrated evidence for functional association of \( I_{Na} \) and \( Na_{a} \) currents through coupling between \( Na_{a} \) and KChIP2, leading to co-regulation in normal circumstances [40]. It might be possible that disruption to the coupling mechanism could explain the findings of differential K\(^+\) channel behaviour in the \( Scn5a+/- \) hearts. The heterogeneities in \( I_{Ko} \) could thus both modify AP repolarization and also influence the reduction of Na\(^+\) current in \( Scn5a+/- \) hearts to a different extent in the RV than the LV. This could explain the protective effects of quinidine in reducing the spatial and temporal electrophysiological heterogeneities and incidence of arrhythmias in BrS [41].

The main limitations of our study come from the differences between mouse and human hearts in the
repolarization phase of the AP, with differences in the range of K⁺ currents displayed. Mouse hearts also use less L-type Ca²⁺ channel current I\textsubscript{Ca,L} with the mouse cardiac AP lacking a plateau phase. The technical difficulty in separating epicardial and endocardial layers meant that our study was limited to a comparison of LV and RV, and we were thus unable to compare transmural differences. Furthermore, we acknowledge that the myocyte isolation procedure is known to cause cell damage, which may affect ionic current properties, including the number of channels and the peak conductance; however, our methodology in the perfusion technique and in the criteria for selection of cells for inclusion in the study aimed to minimize these factors.

Experimental conditions for Na⁺ current measurement were unphysiological, yet chosen for the following reasons. First, voltage-clamp control of the cell membrane was considered. Under physiological conditions, I\textsubscript{Na} is so large and the kinetics are so rapid that successful voltage-clamp control is challenging, even in single-cell preparations [42,43]. For these reasons, we followed protocols that studied I\textsubscript{Na} in a whole-cell recording configuration under reduced experimental temperatures and lower Na⁺ gradients [16,44–46]. As different temperatures were used for I\textsubscript{Na} measurements from those of I\textsubscript{pNa} and I\textsubscript{Ko}, direct comparisons of these currents cannot be made in this paper. Under our conditions, peak I\textsubscript{Na} was 5.4 ± 0.1 nA and 5.3 ± 0.1 nA for WT LV and RV, respectively; these values are similar to others under similar conditions [16,20,47,48]. To enhance voltage-clamp control, large patch pipettes were used. Thus, peak steady-state voltage error resulting from residual series resistance was 2–3 mV, similar to that found in other studies [16,20,47,49]. If experiments demonstrated evidence of inadequate voltage control (e.g. a ‘threshold phenomenon’ near the voltage range for Na⁺ channel activation), data were discarded. The absence of crossover of the current traces as the current magnitude was changed and the shape of the I–V relationship remaining unchanged for difference peak current values further suggest adequate voltage control [16,17]. Only around the reversal potential did we find that the shape of the I–V relationship differed slightly between curves, with the WT RV trace giving a slightly more negative reversal potential than would be expected from the calculated Nernst potential of ~40 mV. Thus, the clamp voltage was properly controlled in our dialysed cell experiments, and it is unlikely that our findings of reduced I\textsubscript{Na} result from clamp conditions.

The Na⁺ flux recorded may have been affected by voltage-dependent blockage by intracellular Mg²⁺, although this appears to be significant only at high Mg²⁺ concentrations and for outward currents [50,51]. Ca²⁺ flux has also been postulated to occur through Na⁺ channels under β-adrenergic stimulation in a process dubbed ‘slip-mode conductance’ [52], although other groups have found conflicting results [53–55]. While classical studies have shown little affinity of Ca²⁺ or Mg²⁺ for the pore site of the channel [56], the use of low external Na⁺ concentrations may increase possible blockade of the Na⁺ channel by either ion.

While decreased membrane excitability owing to reduced Na⁺ channel function may cause slowed conduction in itself, there is also growing evidence that it may also lead to conduction slowing through the generation of fibrosis [10]. As several previous studies have examined this phenomenon in the Scn5a+/− mouse, in this study, we chose to focus on the possible electrophysiological mechanisms underlying the arrhythmias; however, we acknowledge the potential roles of fibrosis in both slowing conduction and also in promoting and maintaining differences in repolarization timing between myocardial layers. Thus, both structural changes and differences in ion channel densities and kinetics could together create optimal pro-arrhythmic conditions.

Changes both in Scn5a+/− mice and in BrS patients could also involve altered expression of a range of other genes and ion channels. Right ventricular outflow tract (RVOT) origins of arrhythmias might also involve slow conducting tissue whose AP upstroke is I\textsubscript{Ca,L}-dependent [57]. Ventricular mRNA expression profiles in studies of human BrS associated with Na⁺ channelopathy have suggested remodelled Ca²⁺ in addition to Na⁺ and K⁺ channel expression [58]. Furthermore, Ca²⁺/K⁺-dependent mechanisms are known to play a role in Na⁺,1,5 regulation [59]. Future work might then explore the possibility that altered current densities and expression levels of Ca²⁺ channels might also contribute to arrhythmogenesis in Scn5a+/− hearts.

The cellular and molecular experiments described here correlate with work performed in the whole heart, especially mapping experiments that have localized both conduction slowing and heterogeneity of repolarization, and subsequent initiation of arrhythmias through reentrant phenomena, to the RV [8]. Computer simulations could provide a potential means to further explore the pro-arrhythmic consequences of RV/LV differences in ionic currents, particularly employing whole heart models [60–63]. However, this would also entail obtaining further independent information on detailed conduction pathway morphology, including extents of fibrosis and cell–cell conductance information in Scn5a+/− hearts. While groups have previously used whole heart models of WT mouse hearts [60], it has been shown previously that Scn5a+/− hearts exhibit heterogeneous fibrosis [2], along with heterogeneous expression of connexin 43 and upregulation of hypertrophic markers including β-MHC and skeletal α-actin [64].

In conclusion, Scn5a+/− mice demonstrate a relative upregulation of Scn5a expression in the LV compared with RV. The specifically reduced RV expression of Na⁺ channels leads to smaller I\textsubscript{Na}, resulting in slowed conduction, and smaller I\textsubscript{pNa} which, in combination with increased I\textsubscript{Ko} results in shorter AP durations and greater heterogeneity of repolarization, thus suggesting that arrhythmogenesis may be initiated by both abnormal depolarization and repolarization in the RV of Scn5a+/− hearts.

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