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Mechanism of Sinus Bradycardia in Carriers of the A414G Mutation in the *HCN4* Gene

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Abstract

Heterozygous carriers of the A414G mutation in the HCN4 gene, which encodes the HCN4 protein, show moderate to severe sinus bradycardia. Tetramers of HCN4 subunits constitute the ion channels that conduct the cardiac hyperpolarization-activated 'funny current' (I_f), which plays an important modulating role in the pacemaker activity of sinus node cells.

We assessed the mechanism by which the A414G mutation in HCN4 causes sinus bradycardia. We carried out voltage clamp experiments on HCN4 channels expressed in Chinese hamster ovary (CHO) cells and incorporated the experimentally observed mutation-induced changes in I_f into the Fabbri-Severi model of a single human sinus node cell.

In the Fabbri-Severi model, the experimentally observed effects on I_f increased the cycle length from 813 to 1004 ms, corresponding with a 19% decrease in beating rate from 74 to 60 beats/min. These mutation effects became more prominent at 10 nM ACh (vagal tone) and in the presence of a hyperpolarizing atrial load.

We conclude that the experimentally identified mutation-induced changes in I_f can explain the clinically observed sinus bradycardia in carriers of the A414G mutation in the HCN4 gene.

1. Introduction

The hyperpolarization-activated 'funny current' (I_f), also known as 'pacemaker current', is a key player in sinoatrial node (SAN) pacemaker activity. This (mainly) inward current of mixed ionic nature is a determinant of the spontaneous depolarization that underlies SAN pacemaker activity and thus a modulator of pacing rate [1,2]. The I_f ion channel in the cell membrane is constituted by four hyperpolarization-activated, cyclic-nucleotide-gated (HCN) subunits. The HCN4 protein, which is encoded by the *HCN4* gene, is the dominant HCN isoform

in rabbit and human SAN [3–6].

Since 2003, several loss-of-function mutations in the *HCN4* gene have been associated with human sinus bradycardia [7,8]. We reviewed these mutations in 2014 [9] and, more recently, in 2015 [10]. Around that time, Milano et al. [11] and Schweizer et al. [12] were the first to report on an intriguing association between loss-of-function mutations in *HCN4* and left ventricular non-compaction (LVNC). Milano et al. [11] observed such non-compaction for the G482R, Y481H, and A414G missense mutations and Schweizer et al. [12] for the G482R and P883R missense mutations, and the 695X truncation mutation [12]. Each of these loss-of-function mutations is also associated with sinus bradycardia.

In the present study, we assessed the mechanism by which the aforementioned A414G mutation in *HCN4* causes sinus bradycardia. To this end, we first carried out voltage clamp experiments on wild-type and heterozygous mutant HCN4 channels expressed in Chinese hamster ovary (CHO) cells. Next, we incorporated the experimentally identified mutation-induced changes in the kinetics of the HCN4 channels into the comprehensive mathematical model of a single human SAN pacemaker cell that was developed by Fabbri et al. [13] (Fabbri-Severi model). This model allows different levels of autonomic tone through the simulated administration of acetylcholine (ACh; vagal tone) or isoprenaline (Iso; β -adrenergic tone). Computer simulations were carried out without autonomic tone, in presence of 10 nM ACh to test vagal effects, and in the absence or presence of an additional outward current with a conductance of 0.4–0.8 pS/pF and a reversal potential of –80 mV to test the effects of sinus node–atrial interactions.

2. Methods

2.1. Patch clamp experiments

The A414G mutation was introduced in wild-type (WT) hHCN4 complementary DNA (cDNA) as described

previously [11]. Chinese hamster ovary (CHO) cells were transiently transfected with lipofectamine using either 2 μg WT cDNA, or, to recapitulate a heterozygous state, 1 μg of WT and 1 μg of mutant HCN4 construct. HCN4 currents were recorded at $36 \pm 0.2^\circ\text{C}$ two days after transfection using the amphotericin-perforated patch-clamp technique. The amplitude, kinetics and reversal potential of the HCN4 current were determined by voltage clamp protocols as described previously [11,14].

The I_f activation curves were fitted with a Boltzmann equation to determine the half-maximum activation voltage ($V_{1/2}$) and slope factor (k). The time constants of I_f (de)activation were obtained by fitting the time course of I_f (de)activation with a monoexponential equation, ignoring the variable initial delay in I_f (de)activation.

2.2. Computer simulations

The heterozygous A414G mutation in *HCN4* was implemented in the CellML code [15] of the Fabbri et al. model [13] by shifting the voltage dependence of the steady-state activation curve and (de)activation time constant of I_f by and -25.7 and -11.3 mV, respectively, based on the data from our patch clamp experiments.

The default Fabbri et al. model [13] has a beating rate of 74 beats/min. This rate was lowered to 58 beats/min (vagal tone) through the simulated administration of 10 nM

ACh. The major effects of the administration of ACh are the activation of the ACh-activated potassium current $I_{K,ACh}$, which is zero in the default model, and the inhibition of I_f through a negative shift in its voltage dependence [13]. The effects of sinus node-atrial interactions were tested through the addition of an ohmic outward current with a conductance of 0.4–0.8 pS/pF and a reversal potential of -80 mV.

The CellML code was edited and run in the Cellular Open Resource (COR) environment [16], version 0.9.31.1409. All simulations were run for a sufficiently long time to reach steady-state behaviour.

3. Results

3.1. Patch clamp experiments

We carried voltage clamp experiments to assess the effects of the heterozygous A414G mutation in *HCN4* on the functional properties of I_f . These experiments revealed a mutation-induced -25.7 mV shift in $V_{1/2}$ of the I_f steady-state activation curve without a significant change in its slope factor (data not shown). This prominent shift was accompanied by a significant -11.3 mV shift in the voltage dependence of the time constant of I_f (de)activation. The mutation did not affect the fully-activated conductance of I_f or its reversal potential.

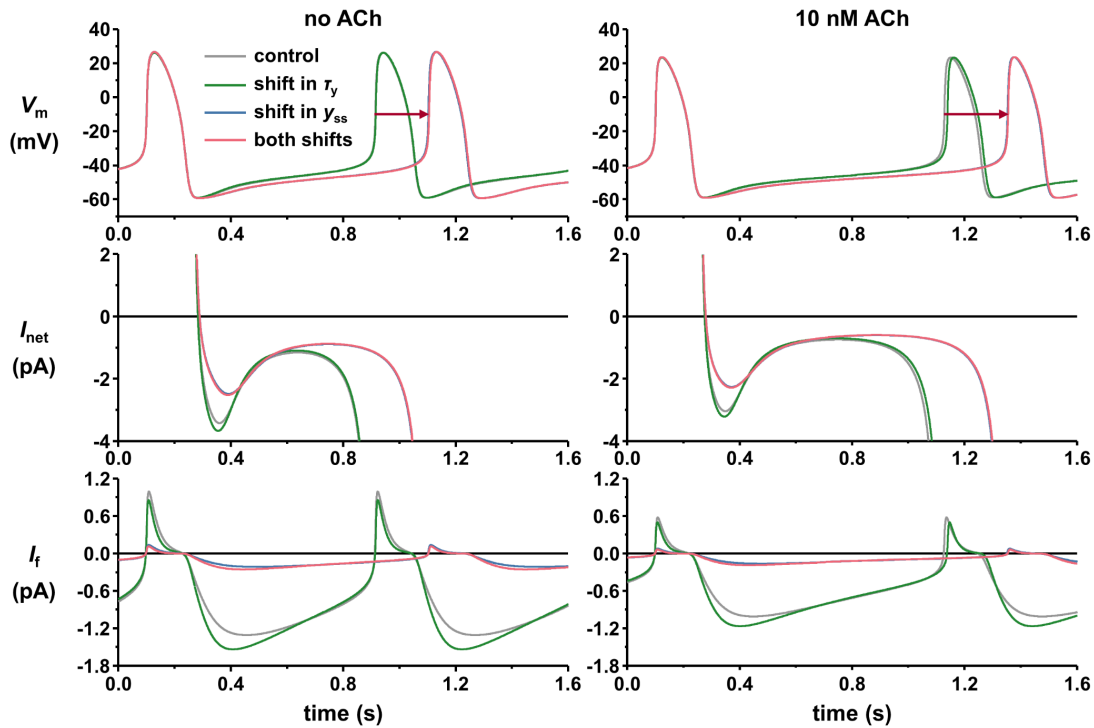


Figure 1. Pacemaker activity in the Fabbri-Severi model of a single human SAN pacemaker cell. Effects of the *HCN4*-A414G mutation-induced -25.7 and -11.3 mV shifts in the steady-state activation y_{ss} and (de)activation time constant τ_y , respectively, of the hyperpolarization-activated ‘funny current’ (I_f) on the membrane potential (V_m , top), net membrane current (I_{net} , middle), and I_f (bottom) under control conditions (no ACh, left) and in the presence of 10 nM ACh (right).

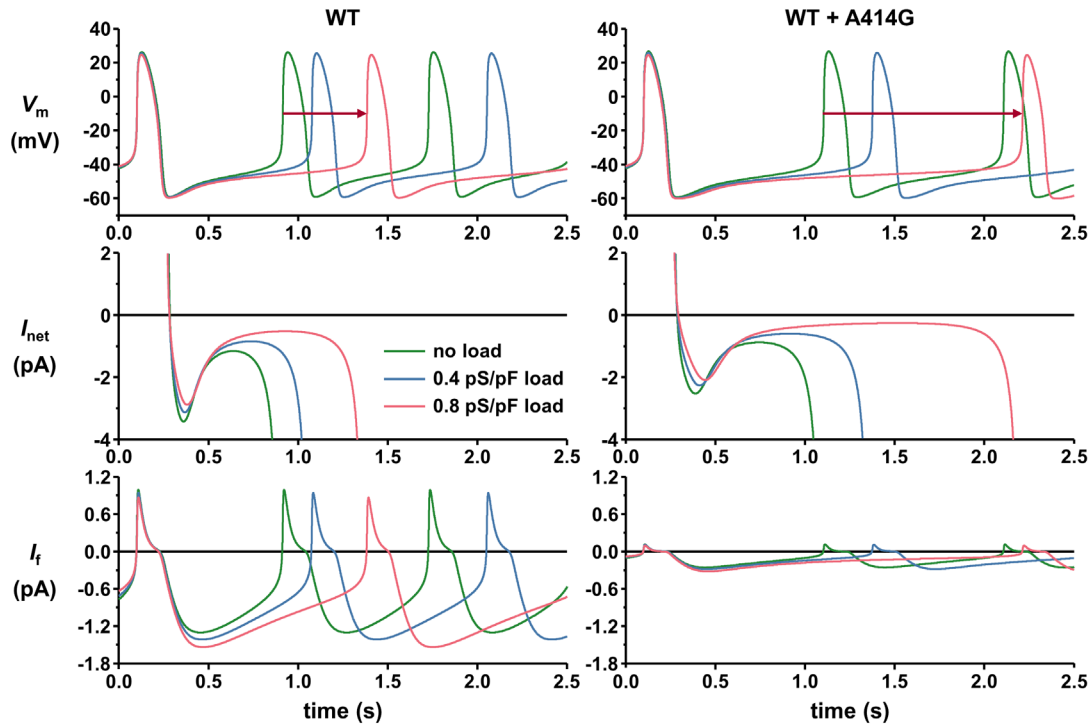


Figure 2. Pacemaker activity in the Fabbri-Severi model of a single human SAN pacemaker cell. Effects of atrial load (i.e., an outward current with a conductance of 0.4–0.8 pS/pF and a reversal potential of -80 mV) on the membrane potential (V_m , top), the net membrane current (I_{net} , middle), and the hyperpolarization-activated ‘funny current’ (I_f , bottom) under wild-type conditions (WT, left) and in the presence of the heterozygous *HCN4*-A414G mutation (WT + A414G, right).

3.2. Computer simulations

Fig. 1 shows the effects of the -25.7 mV shift in the steady-state activation of I_f (y_{ss}) as well as the -11.3 mV shift in its (de)activation time constant (τ_y) on pacemaker activity. The horizontal arrows indicate the mutation-induced increase in cycle length under control conditions (left) and in the presence of 10 nM ACh (right). This increase amounts to 191 and 227 ms, respectively, and is almost completely attributable to the -25.7 mV shift in y_{ss} . The beating rate decreases by 19% (from 74 to 60 beats/min) under control conditions and by 18% (from 58 to 48 beats/min) in the presence of 10 nM ACh.

Fig. 2 shows the effects of atrial load on pacemaker activity under wild-type conditions (left) and in the presence of the heterozygous *HCN4*-A414G mutation (right). In the latter case, the model cell is more sensitive to atrial load, with an increase in cycle length up to 1110 ms (+110%) in case of 0.8 pS/pF load as compared to 470 ms (+58%) under control conditions (horizontal arrows).

The combined effects of atrial load and vagal tone (10 nM ACh) on pacemaker activity are shown in Fig. 3. Under wild-type conditions, an atrial load of 0.4 pS/pF results in a 39% increase in cycle length from 1027 to 1428 ms, whereas an atrial load of 0.6 pS/pF results in a 94%

increase in cycle length to 1988 ms. The effects of atrial load are much more pronounced in the presence of the heterozygous *HCN4*-A414G mutation. An atrial load of 0.4 pS/pF results in a 68% increase in cycle length from 1276 to 2146 ms, whereas an atrial load of 0.6 pS/pF results in a complete loss of pacemaker activity. The beating rate at 0.4 pS/pF amounts to 42 beats/min under wild-type conditions and 28 beats/min in the presence of the heterozygous *HCN4*-A414G mutation

4. Conclusion

We conclude that the experimentally observed changes in the kinetics of I_f can explain the sinus bradycardia that has been identified in carriers of the A414G loss-of-function mutation in *HCN4*.

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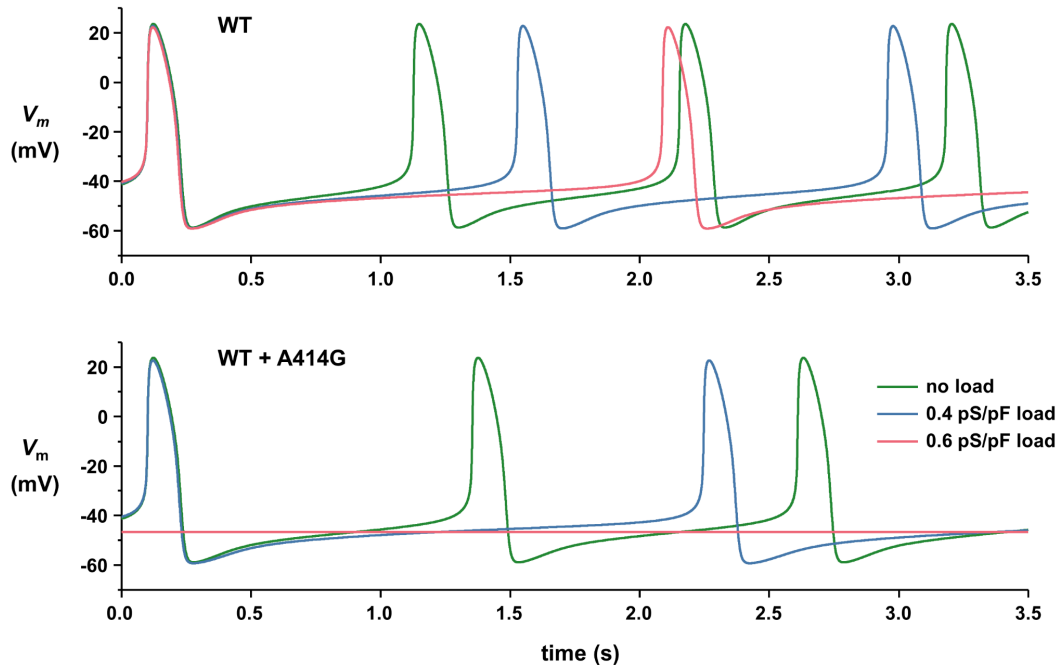


Figure 3. Pacemaker activity in the Fabbri-Severi model of a single human SAN pacemaker cell. Effects of atrial load in presence of vagal tone (10 nM ACh) on the membrane potential (V_m) under control conditions (WT, top) and in the presence of the heterozygous *HCN4*-A414G mutation (WT + A414G, bottom).

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