Persistent human cardiac Na$^+$ currents in stably transfected mammalian cells
Robust expression and distinct open-channel selectivity among Class 1 antiarrhythmics

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Abbreviations: TTX, tetrodotoxin; BTX, batrachotoxin; hNav1.5-CW, human cardiac Na$^+$ channel with L409C/A410W mutations

Introduction

Voltage-gated Na$^+$ channels are responsible for the generation of action potentials in excitable membranes. Upon depolarization neuronal Na$^+$ channels open only once with a brief dwell time of ~0.5 ms.1 These open channels are fast inactivated through the action of an inactivation particle, which rapidly plugs the pore.2 However, it has become increasingly evident that various gain-of-function genetic diseases manifest small but persistent late Na$^+$ currents (generally < 2%) in cardiac myocytes, skeletal muscle and neurons during prolonged depolarization. Among these diseases are congenital long QT-3 syndromes, paramyotonia congenita, potassium-aggravated myotonia and familial primary erythermalgia.3,5 In addition, small persistent Na$^+$ currents have also been found under pathological conditions such as failing heart6 and hypoxia.7 The discoveries of aberrant persistent late sodium currents have attracted much attention recently as a potential target for drug intervention. Therapeutic drugs that eliminate persistent late Na$^+$ currents preferentially may be beneficial for these life-threatening events.

The human cardiac Na$^+$ channel (hNav1.5) is a transmembrane protein,8 which contains four repeated domains (D1-D4) each with six transmembrane segments (S1-S6). Cardiac Na$^+$ channels show longer open times (~1.0 to 2.0 ms), and have a greater likelihood of multiple openings than neuronal counterparts.9 The hNav1.5-ΔKPQ mutant that manifests the long-QT phenotype generates persistent late Na$^+$ currents during prolonged depolarization.10 The site of ΔKPQ deletion occurs at the D3-D4 intracellular linker. Unfortunately, it was rather difficult to study drug binding with these late currents because of their small amplitude.11,12 Large persistent late Na$^+$ currents could be generated by IFM→QQQ triple mutations at the D3-D4

www.landesbioscience.com Channels 263
intracellular linker of the Na+ channel. This IFM motif is identified as a crucial part of the inactivation particle. A diminished inhibitory effect of antiarrhythmic agent lidocaine on the inactivation-deficient hNav1.5-IFM/QQQ mutant channels have been reported. In contrast, flecainide and mexiletine have been found to be potent blockers of small persistent late Na+ currents generated by hNav1.5-ΔKPQ mutant channels. Both of these drugs have been used to treat LQT-3 syndromes.

Attempts to express hNav1.5-IFM/QQQ mutant channels in human embryonic kidney cells (HEK293) cotransfected with an inward rectifier K+ channel (IRK1) yield only partial success. IRK1 channels were introduced to increase the expression of IFM/QQQ mutant Na+ channels by hyperpolarizing the membrane potential. The peak currents in cells expressing the hNav1.5-IFM/QQQ mutant were generally < 2 nA, an amplification of around −80 mV and outward currents appeared around −10 mV. Large non-inactivated Na+ currents were clearly evident during the 80-ms pulse. The sodium conductance was derived from the 80-ms pulse. The sodium conductance was derived from the inward hNav1.5-CW Na+ currents were activated at the threshold value of −41.8 ± 1.4 mV (at voltage where peak current, $E_N$ is the test voltage and $E_{rev}$ is the estimated reversal potential. Normalized $g_N$ values were then plotted against the corresponding voltage (open circles). The plot was fitted with a Boltzmann function. The midpoint voltage ($V_{1/2}$) and slope (k) of the function were −41.8 ± 1.4 mV and 10.2 ± 0.6 mV, respectively (n = 8). The holding potential was set at −140 mV.

Results

Robust expression of hNav1.5-CW channels in a permanent HEK293 cell line. We initially isolated a colony, hNav1.5-CW/8, which expressed a high level of Na+ currents under the whole-cell configuration. However, when this cell line was reestablished from a frozen vial, the expression of Na+ currents dropped precipitously within 3–4 passages. We suspected that a revertant might emerge during the initial selection period. To isolate cells with the mutant Na+ channel clone from the total population of cultured hNav1.5-CW/8 cells, we diluted them -300,000-fold, plated cells into 100-mm cultured dishes, and rescreened for Na+ channel expression as described in the methods section. One rescreened colony was found to remain stable with robust expression of inactivation-deficient hNav1.5-CW Na+ currents for up to 2 mo. This stable cell line (hNav1.5-CW/8–8) was expanded, frozen and stored in a liquid nitrogen tank. Frozen cells were thawed and reestablished in Ti-25 flasks. These cells were kept by continuous passages over a period of 3 mo. G-418 was not needed in the media for the reestablished cell line.

Figure 1A shows superimposed traces of inactivation-deficient hNav1.5-CW Na+ currents. These traces were recorded from a cell reestablished for 2 mo. Under our experimental conditions, inward hNav1.5-CW Na+ currents were activated at the threshold of around −80 mV and outward currents appeared around −10 mV. Large non-inactivated Na+ currents were clearly evident during the 80-ms pulse. The sodium conductance was derived from the peak amplitude, normalized and plotted against membrane potential (Fig. 1B). The graph was fitted using a Boltzmann equation with a $V_{1/2}$ value of −41.8 ± 1.4 mV (at voltage where 50% channels activated, n = 8) and a slope factor of 10.2 ± 0.6 mV (n = 8). These values are comparable to those found in wild-type hNav1.5 channels.

To estimate the level of Na+ channel expression in individual cells we determined its peak amplitude of Na+ current and its capacitance. By and large, these cells expressed 996 ± 173 pA/pF (n = 20) of peak currents at +50 mV. This amount of expression is much higher than that of hNav1.5-IFM/QQQ inactivation-deficient Na+ currents in a permanent HEK293 cell line and at least twice as high as the skeletal muscle rNav1.4-WCW stable...
of currents were non-inactivating during the 4-second pulse. This onset of slow inactivation for inactivation-deficient hNav1.5-CW currents was, however, faster than those found for wild-type hNav1.5 currents. Such phenomenon has been also reported in inactivation-deficient hNav1.5-IFM/QQQ mutant channels and demonstrates that slow inactivation is enhanced in fast inactivation-deficient Na$^+$ channels.

Earlier experiments have demonstrated that batrachotoxin (BTX) could eliminate the slow inactivation of voltage-gated Na$^+$ channels. BTX also significantly alters the activation process as well as changes the ion selectivity of voltage-gated Na$^+$ channels. Our results demonstrated that hNav1.5-L409C/A410W mutant channels could be readily modified by 5 μM BTX. To

![Figure 2](image-url)

**Figure 2.** BTX prevents the slow decay of hNav1.5-CW mutant Na$^+$ currents. (A) A family of superimposed hNav1.5-CW Na$^+$ current traces was recorded with 4-second test pulses, ranging from −120 to +50 mV. Individual current traces at −80, 0 and +50 mV are labeled. The Na$^+$ reversal potential is near −10 mV. (B) A family of superimposed hNav1.5-CW Na$^+$ current traces were recorded with 4-second test pulses ranging from −140 to +50 mV in the presence of 5 μM BTX within the pipette solution. Notice that BTX-modified Na$^+$ currents were maintained throughout the duration of the pulse. Individual current traces at −130, −100, −50, 0 and +50 mV are labeled. Before this record was taken, 400 repetitive 20-ms pulses at +50 mV were applied in a frequency of 2 Hz to facilitate BTX binding with the open channel. The Na$^+$ reversal potential is near −20 mV. Different cells were used in each panel. (C) With 5 μM BTX within the pipette, the Na$^+$ conductance (closed circles) was clearly shifted to the hyperpolarizing direction as compared with that without BTX presence (dashed line as from Fig. 1B). The midpoint voltage ($V_{0.5}$) and slope (k) of the function were −109.7 ± 1.0 mV and 7.6 ± 0.9 mV, respectively (n = 5). Currents were recorded as shown in Figure 2B and the conductance was calculated as described Figure 1B. Holding potential was set at −140 mV.

Evidently, a deficiency in the fast Na$^+$ channel inactivation per se did not impair the stable expression of hNav1.5-CW channels in this permanent HEK293 cell line.

**Slow inactivation of hNav1.5-CW mutant cardiac Na$^+$ channels with and without batrachotoxin.** Human Nav1.5-CW cardiac Na$^+$ currents were visible during prolonged pulses at voltages between −80 mV to +50 mV (Fig. 2A). These persistent late hNav1.5-CW Na$^+$ currents decreased slowly with two exponential decay constants, likely because of the slow inactivation of the hNav1.5-CW mutant channel. The majority of current decay followed a slow time constant of 1.64 ± 0.22 sec at +50 mV (68 ± 3%, n = 6). The remaining decreased with a faster time constant of 129 ± 8 ms (25 ± 2%, n = 6) and about 7 ± 1% (n = 6) of currents were non-inactivating during the 4-second pulse. This onset of slow inactivation for inactivation-deficient hNav1.5-CW currents was, however, faster than those found for wild-type hNav1.5 currents. Such phenomenon has been also reported in inactivation-deficient hNav1.5-IFM/QQQ mutant channels and demonstrates that slow inactivation is enhanced in fast inactivation-deficient Na$^+$ channels.

Earlier experiments have demonstrated that batrachotoxin (BTX) could eliminate the slow inactivation of voltage-gated Na$^+$ channels. BTX also significantly alters the activation process as well as changes the ion selectivity of voltage-gated Na$^+$ channels. Our results demonstrated that hNav1.5-L409C/A410W mutant channels could be readily modified by 5 μM BTX. To
initiate BTX binding, we first applied repetitive pulses (a total of 400 at +50 mV/24 ms) at a frequency of 1 Hz. Figure 2B shows BTX-modified hNav1.5-CW Na+ current traces from −140 to +50 mV at a slow time frame. Inward Na+ currents appeared around −130 mV (Fig. 2B); after activation, these Na+ currents remained during 4-second depolarization with minimal decay. Because of the hyperpolarizing shift in activation by BTX, sizable inward Na+ currents could be observed between −120 to −30 mV (Fig. 2B), whereas only smaller inward Na+ currents were recorded between −70 to −20 mV in the absence of BTX (Fig. 2A). In fact, BTX alters the V50 value for activation by −58 mV (from −41.8 mV to −109.7 mV; Fig. 2C, solid vs. dotted line) and eliminates most of the slow inactivation of these inactivation-deficient hNav1.5-L409C/A410W mutant channels (Fig. 2B). The reversal potential for the BTX-modified hNav1.5-CW Na+ channel was leftward shifted, which reversed around −20 mV, probably due to its reduced selectivity for Na+ ions23 and/or Na+ ion loading. This leftward-shift phenomenon of the reversal potential induced by BTX is commonly found in the literature.

Block of hNav1.5-CW mutant Na+ channels by external TTX at the μM range. We have previously shown that TTX blocked skeletal muscle inactivation-deficient hNav1.4-CW Na+ channels potently (IC50 = 15.1 nM).24 In comparison, TTX block the hNav1.5-CW mutant channels at μM range. Figure 3A shows the superimposed current traces before and after the external application of TTX at various concentrations. The dose-response curve Figure 3B (open circle) shows the IC50 for TTX block is estimated 2.32 ± 0.02 μM (Hill coefficient = 0.98 ± 0.01; n = 6) for the peak Na+ current block (Fig. 3B, open circle), demonstrating that cardiac Na+ channel remains relatively resistant to TTX block. This TTX potency of hNav1.5-CW is about 160-fold weaker than its skeletal muscle hNav1.4-CW counterpart. Our estimated IC50 value is nearly identical to that measured in wild-type hNav1.5 expressed in Xenopus oocytes (tonic block; 2.2 μM) but lower than 7.4 μM for hNav1.5-IFM/QQQ inactivation-deficient mutant channels.25 Furthermore, we found that there was no indication of a TTX-induced time-dependent block as shown in Figure 3B (closed circles). Maintained currents were also blocked by TTX during 50-ms depolarization with an IC50 for TTX block of 2.07 ± 0.04 μM (Hill coefficient = 0.94 ± 0.02; n = 6). The difference in IC50 values for peak and persistent late Na+ currents, however, is statistically significant (with a ratio of 1.12; p < 0.05).

Open-channel block by Class 1c antiarrhythmic drugs flecainide and propafenone during depolarization. Earlier studies indicate that persistent late Nav1.5 Na+ currents could be sensitive to block by Class 1c antiarrhythmic drugs, such as flecainide and propafenone.12,26 We tested these Na+ channel blockers at their therapeutic concentrations to see whether they also effectively blocked hNav1.5-CW Na+ currents in our permanent HEK293 cell line. We found that late hNav1.5-CW currents were far more sensitive to the block of flecainide and propafenone at 0.1 to 10 μM than were peak Na+ currents. Consequently, a strong time-dependent block of hNav1.5-CW Na+ currents by these Class 1c antiarrhythmic drugs appeared during the 100-ms depolarizing pulse. Figure 4A and B show the time-dependent block induced by flecainide and propafenone, respectively. These drugs attained their steady-state block of late hNav1.5-CW Na+ currents at +50 mV within 1.5–5 min. The dose-response curves of the peak and maintained late hNav1.5-CW Na+ currents for flecainide and propafenone are shown in Figure 5A and B, respectively. These curves were fitted by a Hill equation, and the estimated 50% inhibitory concentrations (IC50 values) and Hill coefficient for the resting and the open states of Nav1.5-CW Na+ channels are listed in Table 1. We define the resting-channel block as the block of the peak Na+ currents and the open-channel block as

Figure 3. TTX blocks hNav1.5-CW Na+ channels. (A) Superimposed Na+ current traces were recorded in the absence and in the presence of TTX at 1, 3 and 10 μM. Currents were evoked by a 50-ms test pulse to +50 mV every 30 seconds. A steady-state block at each concentration was reached before application of the next solution. The block of Na+ currents by TTX could be completely reversed by the wash of the bath solution. (B) The dose-response curves were constructed using the data set as shown in (A). The peak or late Na+ currents were measured, normalized to the amplitude of the control, and plotted against the TTX concentration. Solid line represents a fit to the data with the Hill equation. The estimated IC50 values ± standard error (Hill coefficient ± S.E.) for peak-current block and the late-current block were 2.32 ± 0.02 μM (0.98 ± 0.01) (n = 6) and 2.07 ± 0.04 μM (0.94 ± 0.02) (n = 6), respectively.

Figure 4. Class 1c antiarrhythmic drugs flecainide and propafenone block hNav1.5-CW Na+ currents. (A) Superimposed Na+ currents before and after the wash of the bath solution. The block of Na+ currents by external application of the next solution. The block of Na+ currents by TTX could be completely reversed by the wash of the bath solution. (B) The dose-response curves were constructed using the data set as shown in (A). The peak or late Na+ currents were measured, normalized to the amplitude of the control, and plotted against the TTX concentration. Solid line represents a fit to the data with the Hill equation. The estimated IC50 values ± standard error (Hill coefficient ± S.E.) for peak-current block and the late-current block were 2.32 ± 0.02 μM (0.98 ± 0.01) (n = 6) and 2.07 ± 0.04 μM (0.94 ± 0.02) (n = 6), respectively.
block at the peak amplitude measurement may also contribute to the deviation of the Hill coefficient for flecainide from the unity (0.52 ± 0.04, n = 6; Table 1). Our results nonetheless demonstrate clearly that (1) persistent late hNav1.5-CW Na\(^{+}\) currents in stably transfected HEK293 cells retain their drug sensitivity toward Class 1c antiarrhythmics at clinic relevant concentrations, and

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**Figure 4.** Block of hNav1.5-CW Na\(^{+}\) channels by Class 1c antiarrhythmic drugs. Superimposed hNav1.5-CW Na\(^{+}\) currents were recorded before and after the application of flecainide (A) or propafenone (B) at concentrations of 0.1, 0.3, 1.0, 3.0 and 10 \(\mu\)M. Currents of hNav1.5-CW Na\(^{+}\) channels were evoked every 30 seconds by a 100-ms test pulse at +50 mV. Drugs were applied externally in the external solution, and the currents were recorded after the block reached its steady-state, usually within 5 min. Different cells were used in (A and B). Holding potential was set at −140 mV. Traces of the control (0 \(\mu\)M), 0.3 and 3.0 \(\mu\)M drug concentrations were labeled.

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**Figure 5.** Dose-response curves for Class 1c antiarrhythmic drugs. (A) Dose-response curves for open-channel block and resting-channel block of flecainide were constructed using the data set as described in Figure 4A. For resting-channel block (open circles), we measured the peak amplitude of Na\(^{+}\) currents at various flecainide concentrations, normalized to the peak amplitude of the control, and plotted against drug concentration. Solid lines represent fits to the data with the Hill equation, \(y = 1/[1 + (x/IC_{50})^n]\), where \(IC_{50}\) is the 50% inhibitory concentration and \(n\) is the Hill coefficient. The \(IC_{50}\) value (mean ± SEM) and the Hill coefficient (mean ± SEM) were estimated 50.8 ± 11.5 \(\mu\)M and 0.52 ± 0.04, respectively (n = 8), from the fitted curve. For open-channel block (filled circles), we measured the late current amplitude near the end of the 100-ms pulse at various flecainide concentrations, normalized to the late amplitude of the control, and plotted against drug concentration. The \(IC_{50}\) value (mean ± SEM) and the Hill coefficient (mean ± SEM) were estimated 0.63 ± 0.06 \(\mu\)M and 0.95 ± 0.08, respectively (n = 8). (B) Dose-response curves for propafenone were similarly constructed as described above. For resting-channel block (open circles), the \(IC_{50}\) value (mean ± SEM) and the Hill coefficient (mean ± SEM) were estimated 12.0 ± 0.7 \(\mu\)M and 0.98 ± 0.05, respectively (n = 8). For open-channel block (filled circles), the \(IC_{50}\) value (mean ± SEM) and the Hill coefficient (mean ± SEM) were estimated 0.51 ± 0.01 \(\mu\)M and 1.04 ± 0.02, respectively (n = 8).

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the block of the persistent late Na\(^{+}\) currents near the end of the 100-ms pulse at +50 mV. However, the \(IC_{50}\) values for the resting-channel block shown in Figure 5 are likely higher since a fraction of the open-channel block will occur inevitably at the time when the current reaches its peak amplitude, particularly at high drug concentrations. Such contamination of the resting-channel block at the peak amplitude measurement may also contribute to the deviation of the Hill coefficient for flecainide from the unity (0.52 ± 0.04, n = 6; Table 1). Our results nonetheless demonstrate clearly that (1) persistent late hNav1.5-CW Na\(^{+}\) currents in stably transfected HEK293 cells retain their drug sensitivity toward Class 1c antiarrhythmics at clinic relevant concentrations, and
Table 1. Estimates of the 50% inhibitory concentration (IC$_{50}$) and Hill coefficient of antiarrhythmics for resting and open states of human Nav1.5-CW mutant channels

| Antiarrhythmic drugs | IC$_{50}$ value (μM) and Hill coefficient for resting state | IC$_{50}$ value (μM) and Hill coefficient for open state | Ratio of IC$_{50}$ (resting/open) |
|----------------------|----------------------------------------------------------|----------------------------------------------------------|---------------------------------|
| Flecainide (Class 1c) | 50.8 ± 11.5 μM (0.52 ± 0.04) | 0.63 ± 0.06 μM (0.95 ± 0.08) | 80.6 |
| Propafenone (Class 1c) | 12.0 ± 0.7 μM (0.98 ± 0.05) | 0.51 ± 0.01 μM (1.04 ± 0.02) | 23.5 |
| Lidocaine (Class 1b) | 606 ± 15 μM (0.71 ± 0.01) | 35.3 ± 2.7 μM (0.82 ± 0.05) | 17.2 |
| Mexiletine (Class 1b) | 220 ± 54 μM (0.76 ± 0.16) | 14.9 ± 0.9 μM (0.90 ± 0.05) | 14.8 |
| Quinidine (Class 1a) | 72.7 ± 2.9 μM (1.38 ± 0.07) | 8.6 ± 0.4 μM (1.19 ± 0.07) | 8.5 |
| Disopyramide (Class 1a) | 158 ± 11 μM (1.02 ± 0.08) | 52.5 ± 2.8 μM (1.05 ± 0.05) | 3.0 |

IC$_{50}$ values for resting and open states of hNav1.5-CW channels were estimated as described in Figure 5. Mean and SE of IC$_{50}$ and Hill coefficient (in parenthesis) are listed (each with n = 6–8).

(2) the open hNav1.5-CW Na$^+$ channel interacts preferentially with flecainide and propafenone, at least 20-fold more so than does its resting counterpart.

It is noteworthy that data shown in Figure 4 could be further analyzed to extract the on- and off-rate constants. For example, the time-dependent block at various drug concentrations can be first normalized with the control record in the absence of drug (for removing the slow decay component in the control trace); then the corrected current trace can be fitted with a single exponential function to obtain the time constant ($\tau$). The plot of $1/\tau$ vs. drug concentration can be best fitted with a linear function, which yielded a slope and a y intercept. The values of the slope and the y intercept are equal to the on- and off-rate constants, respectively. Details of such analyses have been described previously. We were able to obtain the on- and off-rate constant for propafenone of 44.2 ± 2.8 μM$^{-1}$s$^{-1}$ and 14.7 ± 1.5 sec$^{-1}$, respectively (n = 6). The calculated dissociation constant ($K_d$ = off-rate/on-rate or 14.7/44.2) yielded 0.33 μM, a value comparable to the estimated IC$_{50}$ (Table 1). Since the slow inactivation is eliminated by normalization, the time-dependent block induced by the Class 1 antiarrhythmic is likely to reflect solely the open-channel block. Similar analyses were employed for flecainide, which yielded an on- and off-rate constants of 14.6 ± 1.2 μM$^{-1}$s$^{-1}$ and 6.6 ± 0.9 sec$^{-1}$, respectively (n = 6), and the $K_d$ value of 0.45 μM. Thus, this cell line is well-suited for detailed kinetic analyses of interactions between drug and open Na$^+$ channels.

Open-channel block by Class 1b antiarrhythmic drugs lidocaine and mexiletine during depolarization. Earlier studies have indicated that the block of the inactivated state of cardiac Na$^+$ channels by lidocaine and mexiletine plays a dominant role in their therapeutic action. However, both of these Class 1b antiarrhythmics also blocked the persistent late hNav1.5-CW Na$^+$ currents effectively. We observed the strong time-dependent block of the late Na$^+$ currents with lidocaine or mexiletine at their clinic relevant concentrations (Fig. 6A and B), similar to those found in Figure 4 for Class 1c antiarrhythmics. This time-dependent block phenotype has previously been well characterized in inactivation-deficient muscle Na$^+$ channels. Figure 7A and B show the dose-response curves of lidocaine and mexiletine for the block of peak and late Na$^+$ currents generated by hNav1.5-CW channels, respectively. Table 1 lists the IC$_{50}$ of lidocaine and mexiletine for resting state (as measured by the peak current block) and open state (as measured by the late current block). The difference of the IC$_{50}$ values between the resting and open-channel block is about 15-fold. These results suggest that, in addition to the dominant inactivated-channel block induced by lidocaine and mexiletine at their therapeutic concentrations, the open-channel block of the cardiac Na$^+$ channel could also play a significant role in their antiarrhythmic action.

Block of persistent late hNav1.5-CW Na$^+$ currents by Class 1a antiarrhythmics disopyramide and quinidine. Grant et al. (2000) reported a lack of time-dependent block of hNav1.5-IFM/QQQ inactivation-dependent Na$^+$ currents by disopyramide at 200 μM. We thought to corroborate such a finding using the stably transfected cell line as an assay to distinguish the action of disopyramide from other Class 1 antiarrhythmic drugs. Figure 8A shows the superimposed traces of hNav1.5-CW Na$^+$ currents before and after external disopyramide application at various concentrations. We found that a stronger block of the late Na$^+$ currents than that of the peak Na$^+$ currents was still present, albeit less severe than that found for Class 1c and 1b antiarrhythmic drugs. The estimated difference in IC$_{50}$ values between the block of peak and the late Na$^+$ currents for disopyramide was about 3-fold (Fig. 9A), which is much less than those of the Class 1c and 1b antiarrhythmics (Table 1). Our results thus demonstrate that the phenotype for disopyramide block in hNav1.5-CW and hNav1.5-IFM/QQQ mutant channels is quite...
(80.6) > propafenone (23.5) > lidocaine (17.2) > mexiletine (14.8) > quinidine (8.5) > disopyramide (3.0). The large disparity in this ratio indicates that Class 1c and 1b antiarrhythmics target the persistent late Na$^+$ currents rather selectively whereas the Class 1a drugs show only a modest selectivity toward late Na$^+$ currents.

**Discussion**

Miniature persistent Na$^+$ currents in cardiomyocytes can cause arrhythmia and sudden death.5,5 Because of their minute size
it has been difficult to study these aberrant late currents and to identify therapeutic drugs that may target them selectively. Details of mutant channel behavior, which determine drug interactions and the functional effects of discrete mutations, are likely to depend on both the channel isoform and the species. This report describes for the first time an HEK293 cell line stably transfected with an inactivation-deficient mutant Na⁺ channel clone that can generate large persistent late human cardiac Na⁺ currents. Pharmacological profiles and gating properties confirm the identity of the hNav1.5-CW mutant Na⁺ channel. With such a cell line available we can also address an open question regarding the open-channel vs. resting-channel selectivity among Class 1 antiarrhythmics.

**Robust expression of hNav1.5-L409C/A410W mutant Na⁺ channels in a stable HEK293 cell line.** Based on their attempts to establish stably transfected Hek293 cell lines expressing inactivation-deficient cardiac hNav1.5-IFM/QQQ Na⁺ channels, Grant et al. suggested that the overload of Na⁺ ions from impetuous openings could inhibit hNav1.5-IFM/QQQ mutant channel expression of these cells. After numerous trials we were uncertain...
whether robust expression of hNav1.5-CW Na⁺ channels could be attained in a permanent HEK293 cell line. Particularly, we witnessed the rapid loss of inactivation-deficient Na⁺ currents in cells isolated from a colony with a high level of Na⁺ channel expression. This observation could be interpreted that the persistent flow of inward Na⁺ current may be detrimental to the cells. However, with a second colony selection, these isolated cells remain stable in expressing hNav1.5-CW mutant channels. We therefore conclude that robust expression of hNav1.5-CW Na⁺ channels can be achieved in HEK293 cells despite the reported low expression in the hNav1.5-IFM/QQQ inactivation-deficient mutant.

The availability of a permanent HEK293 cell line expressing robust mutant hNav1.5-CW Na⁺ channels has a number of advantages. First, the cell line will make it easier for routine exploitations of inactivation-deficient Na⁺ channels, including biophysical and pharmacological studies of persistent late cardiac hNav1.5-CW Na⁺ currents. Second, the permanent cell line may be applicable to identify potent open-channel selective Class 1 antiarrhythmic drugs using an automated parallel patch-clamp system. There is one disadvantage of using a permanent hNav1.5-CW mutant cell line as the potential drug-binding site may be changed unintentionally by mutations. Such possibility should be taken into consideration when the drug affinity is significantly different from that found in the literature.

**TTX and BTX receptors in hNav1.5-L409C/A410W mutant Na⁺ channels.** Both the TTX receptor and the BTX receptor within the Na⁺ channel have been well characterized, and their locations are mapped along the Na⁺ permeation pathway. To check if the binding sites for TTX and BTX within Na⁺ channels are altered, we studied the effects of these toxins in cells expressing hNav1.5-L409C/A410W cardiac mutant Na⁺ channels. Our results demonstrated that the binding sites for TTX and BTX in hNav1.5-CW Na⁺ channels appeared unchanged (Fig. 2; Fig. 3), as their phenotypes induced by these toxins were similar to those found in wild-type counterparts when treated with these ligands. An unchanged binding site for BTX indicates that the BTX receptor within the inner cavity of hNav1.5-CW Na⁺ channels is comparable to that of the wild-type hNav1.5 channels, whereas an unchanged binding site for TTX suggests that the TTX receptor at the external entryway is also near normal.

It is noteworthy that the cardiac hNav1.5 isoform is relatively TTX insensitive, requiring a high concentration of TTX to block this Na⁺ channel. We found that TTX blocked both peak and persistent late hNav1.5-CW Na⁺ currents without a significant time-dependent block during a 50-ms pulse, although it appeared slightly more potent in the block of the late rather than the peak currents (Fig. 3). A clear time-dependent block was found in TTX block of hNav1.5-QQQ mutant during a long 4 sec depolarizing pulse. These authors suggested that the extra block of the hNav1.5-QQQ channel found during depolarization is due to a higher-affinity TTX site, which is modulated by the activation gating of the sodium channel. It remains unknown whether the slow inactivation plays a role in this enhanced block since the slow inactivation is clearly evident in hNav1.5-CW mutant channels during a prolonged pulse (Fig. 2A).

**Characterization of open-channel blockers using the stable cell line expressing large inactivation-deficient hNav1.5-CW sodium currents.** Class 1 antiarrhythmic drugs target voltage-gated hNav1.5 Na⁺ channels for their therapeutic action. These drugs are subgrouped as 1a, 1b and 1c, roughly dependent upon their rates of recovery from drug-induced block under physiological conditions. Unlike the TTX block which shows little time-dependent block of inactivation-deficient Na⁺ currents during depolarization (Fig. 3), we found that antiarrhythmics, including Class 1c (flecainide, propafenone), Class 1b (mexiletine, lidocaine) and Class 1a (quinidine and disopyramide), blocked persistent late hNav1.5-CW mutant Na⁺ currents far more effectively than they blocked peak Na⁺ currents during the depolarizing pulse (Fig. 4; Fig. 6; Fig. 8; Table 1). The Hill coefficient values for the open-channel block are close to the unity ranging from 0.82 for lidocaine to 1.19 for quinidine, suggesting that one drug molecule blocks one open channel. This result is consistent with the finding that there is one local anesthetic/antiarrhythmic receptor within the inner cavity of the Na⁺ channel with drug-sensing residues situated at multiple S6 segments. The Hill coefficient values for the resting state, however, deviate from the unity considerably ranging from 0.52 for flecainide to 1.38 for quinidine. One explanation for this deviation is because the resting block is measured as the block of peak Na⁺ currents. This type of measurement is undoubtedly tainted by the open block of Na⁺ channels during the channel activation. An additional possibility is that there are multiple resting states and each may have a different affinity toward the antiarrhythmic drug.

The selectivity ratio of IC₅₀ between the resting-channel block and the open-channel block among Class 1 antiarrhythmic drugs differs significantly, ranging from 80.6 for flecainide to 3.0 for disopyramide (Table 1). Flecainide can therefore be viewed as a genuine open-channel selective blocker whereas disopyramide cannot. For disopyramide, the resting, open and inactivated blocks of Na⁺ channels all likely play an important role in its efficacy in vivo. The reason behind this considerable difference in the selectivity ratio of IC₅₀ between the resting-channel and the open-channel block among Class 1 antiarrhythmic drugs is unclear. It is possible that flecainide and propafenone may have a better induced-fit with their receptor in the open state of the hNav1.5-CW channel than with their counterpart in the resting state because of the dynamic movement of S6 segments during Na⁺ channel activation.

**Physiological significance of the open-channel block by Class 1 antiarrhythmic drugs.** Several studies have given two contradictory views concerning the therapeutic role of the open-channel block by lidocaine. First, QX-314 (a quaternary ammonium derivative of lidocaine) fails to inhibit the persistent late Na⁺ currents in pronase-treated squid axons. Second, the open-state binding affinity for lidocaine or cocaine is 3- to 10-fold lower than the inactivated-state binding affinity in hNav1.5-IFM/QQQ channels. These results suggest that the open Na⁺ channel is not blocked preferentially by lidocaine and therefore the inactivated channel plays the most important role in QX-314/lidocaine/cocaine block of Na⁺ channels, in agreement with the modulated receptor hypothesis. Additional results of a weak
open-channel block by flecainide, RAD-243, and disopyramide in hNav1.5-IFM/QQQ mutant channels provide further support for this notion.

In contrast to these findings, various local anesthetics and Class 1 antiarrhythmics preferentially block persistent late Na+ currents created by various conditions. Included are (1) choline-T-treated axons, (2) gain-of-function genetic diseases that cause LQT-3 symptoms, (3) hypoxia/failing cardiac myocytes and (4) inactivation-deficient S6 mutants. These conflicting results lead to an alternative view, in that, to the inactivated state of the Na+ channel, the open state also interacts preferentially with lidocaine, mexiletine and flecainide and that such open-channel block is important therapeutically in vivo when persistent late Na+ currents are present.

It is evident that aberrant persistent late Na+ currents generated via chemical/enzymatic modifications, gain-of-function genetic diseases, site-directed mutagenesis or pathological environment are the result of structural alterations of voltage-gated Na+ channels. Theoretically, such alterations could bring about a decrease or an increase in binding affinities for Class 1 antiarrhythmic drugs. In other words, not all persistent late Na+ currents are created equal in their sensitivity toward Class 1 antiarrhythmic drugs. This possibility would account for the two opposing views described above regarding the therapeutic role of the open-channel block. Unfortunately, it is difficult to measure directly the open-channel block in wild-type Na+ channels because these channels showed a brief open time of ~0.5 to ~2.0 ms. Aside from this uncertainty, the immediate need is to identify drugs that can selectively impede aberrant persistent late Na+ currents in patients, while leaving normal transient Na+ currents intact. The cell line expressing robust human cardiac hNav1.5-CW channels should be applicable as a tool to identify novel Class 1 antiarrhythmic drugs that preferentially silence persistent late cardiac Na+ currents.

Materials and Methods

Selection of HEK293 cells with stably transfected hNav1.5-L409C/A410W cDNA clone. Full-length wild-type hNav1.5 cDNA in pcDNA3 vector was provided by Dr. Augustus Grant (Duke University, School of Medicine). Mutagenesis of hNav1.5-L409C/A410W cDNA was performed as reported previously. The site-directed mutations occurred at position 21–22 of the D1S6 C-terminus. HEK293 cells were purchased from American Type Culture Collection (ATCC) and maintained at 37°C in a 5% CO2 incubator in DMEM (Life Technologies, Inc.) containing 10% fetal bovine serum (HyClone) and 1% penicillin and streptomycin solution (Sigma). HEK293 cells were transfected with the mutant clone by a calcium phosphate precipitation method. One day after transfection, HEK293 cells were treated with 1.0 mg/ml G-418 (Invitrogen, Inc.) in 100-mm culture dishes. About 2 weeks after transfection individual G-418-resistant colonies were isolated using glass cylinders (i.d. = 8 mm). Isolated colonies were trypsinized and replated in 35-mm culture dishes each with a gelatin-treated coverslip. The coverslip was removed from the dish after 2–3 d, and ~5 cells from each coverslip were assayed under the whole-cell configuration. Five positive colonies with inactivation-deficient currents were expanded and frozen individually. One colony initially expressing robust inactivation-deficient sodium currents was later expanded, diluted proportionally to the cell density and subjected to a second colony-selection procedure in 100-mm culture dishes without G-418. The second colony-selection procedure was necessary for this particular colony because the number of cells with the robust expression of inactivation-deficient currents decreased drastically after a few passages. Three out of five colonies were tested positive after the second screening and one remained stable with high expression for up to 2 mo. This colony (identified as hNav1.5-CW/8–8) was expanded and maintained in Ti-25 flasks for studies described here.

Electrophysiology and data acquisition. The whole-cell configuration of a patch-clamp technique was used to study Na+ currents in HEK293 cells at room temperature (22 ± 2°C). Electrode resistance was measured about 0.5 MΩ. Command voltages were elicited with pCLAMP9 software and delivered by Axopatch 200B (Axon Instrument). Cells were held at −140 mV and dialyzed for −10–15 min before current recording. Most of the capacitance and leak currents were cancelled with a patch-clamp device and by P/−4 subtraction. Liquid junction potential was not corrected. Access resistance was about 1 MΩ; series resistance compensation of ~95% typically resulted in voltage errors of ≤ 4 mV at +50 mV. Most current measurements were performed at +50 mV for the outward Na+ currents. Such recordings allowed us to avoid the complication of series resistance artifacts and to minimize inward Na+ ion loading. Conductance was calculated as gNa = INa/(ENa − ENa), where INa is the peak current, ENa is the test voltage and ENa is the estimated reversal potential. Conductance curves (gNa vs. ENa) were fitted by a Boltzmann equation, gNa = gNa,0/[1 + exp((VNa − ENa)/k)], where VNa,0 and k are derived as the midpoint voltage and slope factor of the fit, respectively, ENa is the test potential and gNa,0 is the maximal Na+ conductance. Dose-response curves were fitted with the Hill equation, y = 1/[1 + (x/IC50)ⁿ], where IC50 is derived as the 50% inhibitory concentration and n as the Hill coefficient. Curve fitting was performed by Microcal Origin. A student’s unpaired t-test was used to evaluate estimated parameters (mean ± SE or fitted value ± SE of the fit); P values of < 0.05 were considered statistically significant.

Solutions and chemicals. Flecainide-HCl, propafenone-HCl, lidocaine-HCl, mexiletine-HCl, disopyramide sulfate and quinidine HCl were all purchased from Sigma. Tetrodotoxin (TTX) was purchased from Calbiochem Corp., and batrachotoxin (BTX) was a generous gift from Dr. John Daly. TTX at 1 mM was dissolved in H2O and disopyramide sulfate at 100 mM was dissolved in 20 mM HCl. The remaining drugs were dissolved in dimethylsulfoxide (DMSO) as stock solutions and stored at 4°C. Final drug concentrations were made by serial dilution. The highest DMSO concentration in bath solution was 0.1% except for BTX, which contained 1% DMSO in pipette solution. DMSO at a final concentration of 1% had no effect on Na+ currents. Cells were perfused with an extracellular solution containing (in mM) 65 NaCl, 85 choline-Cl, 2 CaCl2 and 10 HEPES (titrated with tetramethylammonium-OH to pH 7.4).
The pipette (intracellular) solution consisted of (in mM) 100 NaF, 30 NaCl, 10 EGTA and 10 HEPES (titrated with cesium-OH to pH 7.2). The rationale of solution choices is given in the last paragraph.

Disclosure of Potential Conflicts of Interest
No potential conflicts of interest were disclosed.

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The rationale of solution choices is given in the last paragraph.
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