Mechanism of sodium channel Na\textsubscript{v}1.9 potentiation by G-protein signaling

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Tetrodotoxin (TTX)-resistant voltage-gated Na (Na\textsubscript{v}) channels have been implicated in nociception. In particular, Na\textsubscript{v}1.9 contributes to expression of persistent Na current in small diameter, nociceptive sensory neurons in dorsal root ganglia and is required for inflammatory pain sensation. Using ND7/23 cells stably expressing human Na\textsubscript{v}1.9, we elucidated the biophysical mechanisms responsible for potentiation of channel activity by G-protein signaling to better understand the response to inflammatory mediators. Heterologous Na\textsubscript{v}1.9 expression evoked TTX-resistant Na current with peak activation at −40 mV with extensive overlap in voltage dependence of activation and inactivation. Inactivation kinetics were slow and incomplete, giving rise to large persistent Na currents. Single-channel recording demonstrated long openings and correspondingly high open probability (P\textsubscript{o}) accounting for the large persistent current amplitude. Channels exposed to intracellular GTP\textsubscript{S}, a proxy for G-protein signaling, exhibited twofold greater current density, slowing of inactivation, and a depolarizing shift in voltage dependence of inactivation but no change in activation voltage dependence. At the single-channel level, intracellular GTP\textsubscript{S} had no effect on single-channel amplitude but caused an increased mean open time and greater P\textsubscript{o} compared with recordings made in the absence of GTP\textsubscript{S}. We conclude that G-protein activation potentiates human Na\textsubscript{v}1.9 activity by increasing channel open probability and mean open time, causing the larger peak and persistent current, respectively. Our results advance our understanding about the mechanism of Na\textsubscript{v}1.9 potentiation by G-protein signaling during inflammation and provide a cellular platform useful for the discovery of Na\textsubscript{v}1.9 modulators with potential utility in treating inflammatory pain.

INTRODUCTION

Voltage-gated Na (Na\textsubscript{v}) channels are critical for the initiation and propagation of action potentials in excitable tissues, including the brain and peripheral nerves. Two specific isoforms, Na\textsubscript{v}1.8 and Na\textsubscript{v}1.9 (also known as SNS and NaN, respectively), are tetrodotoxin (TTX)-resistant Na\textsubscript{v} channels expressed in the peripheral nervous system (Dib-Hajj et al., 1998; Tate et al., 1998; Akopian et al., 1999; Persson et al., 2010). These channels are also found in the central nervous system (Jeong et al., 2000; Blum et al., 2002; O’Brien et al., 2008). Both Na\textsubscript{v}1.8 and Na\textsubscript{v}1.9 have been implicated in nociception, including neuronal pain signaling triggered by inflammation (Lai et al., 2004).

Na\textsubscript{v}1.8 channels are expressed in retinal amacrine and ganglion cells (O’Brien et al., 2008), small and medium-sized dorsal root ganglion (DRG) neurons, and their nociceptive afferent fibers (Benn et al., 2001). In nociceptive fibers, Na\textsubscript{v}1.8 channels are responsible for slowly inactivating Na currents that contribute to the depolarizing phase of action potentials in C-type small DRG neurons (Renganathan et al., 2001). Na\textsubscript{v}1.9 channels are found in the hippocampus, cortex (Jeong et al., 2000; Blum et al., 2002), photoreceptors and Müller glia (O’Brien et al., 2008), small diameter, nociceptive sensory neurons in DRG (Fang et al., 2002), trigeminal ganglia, and in the intrinsic sensory neurons of the gut (Rugiero et al., 2003; Padilla et al., 2007). Compared with Na\textsubscript{v}1.8 and TTX-sensitive neuronal channels, Na\textsubscript{v}1.9 exhibits unique biophysical properties that include a hyperpolarized voltage-dependent activation, activation and inactivation curves that overlap near the resting membrane potential, slow activation and inactivation kinetics, and a very large persistent current (Cummins et al., 1999; Dib-Hajj et al., 2002; Coste et al., 2004). Persistent current generated by Na\textsubscript{v}1.9 has been proposed to set thresholds for excitability of nociceptive sensory neurons by modulating both the resting potential and responses to subthreshold stimuli (Herzog et al., 2001; Priest et al., 2005; Ostman et al., 2008). A link between Na\textsubscript{v}1.9-associated persistent current and pain sensation was demonstrated in neurons from Na\textsubscript{v}1.9 knockout mice that lack persistent Na current and have greatly reduced inflammatory hyperalgesia (Priest et al., 2005; Amaya et al., 2006).

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Inflammation caused by tissue damage results in pain, reflecting an increase in excitability of the primary afferent neurons innervating the area. Inflammatory agents such as bradykinin, ATP, histamine, prostaglandin E2, and norepinephrine potentiate NaV1.9 current, increasing the excitability of DRG neurons, but these mediators fail to sensitize sensory neurons in NaV1.9-null mice (Maingret et al., 2008a; Ritter et al., 2009). Studies using prostaglandin E2, protein kinase C, or G proteins also have demonstrated a link between inflammatory pathways and NaV1.9-mediated nociceptor excitability (Baker et al., 2003; Baker, 2005). Together, these results indicate that NaV1.9 contributes to the hyperexcitability of nociceptors observed during inflammatory pain. Inflammatory mediators that activate G-protein-dependent signal transduction increase the NaV1.9 Na current (Rush and Waxman, 2004). Moreover, intracellular GTPyS (a hydrolysis-resistant GTP analogue) also potentiates persistent TTX-resistant Na current in wild-type and NaV1.8-null mice sensory neurons (Baker et al., 2003) but does not increase persistent TTX-resistant Na current in small sensory neurons from NaV1.9 knockout mice (Ostman et al., 2008). Discerning the mechanism for GTPyS potentiation of NaV1.9 is important for understanding the encoding of painful sensations mediated by G-protein activation.

Heterologous expression of NaV1.9 has proven difficult, and this has limited opportunities to elucidate functional properties of the channel and to implement small molecule screening assays to discover NaV1.9 modulators. In this study, we used a highly efficient transposon system (Kahlig et al., 2010) to achieve stable expression of human NaV1.9 in heterologous cells. This expression system allowed us to elucidate detailed biophysical properties of the channel, including the single-channel mechanism underlying potentiation by GTPyS. Our work further illustrates a cellular platform useful for the discovery of NaV1.9 modulators that may have utility in treating chronic and inflammatory pain.

MATERIALS AND METHODS

Cell culture
All experiments were conducted using ND7/23 cells (Sigma-Aldrich) grown at 37°C with 5% CO₂ in Dulbecco’s modified Eagle’s medium (DMEM) supplemented with 10% fetal bovine serum (ATLANTA Biologicals), 2 mM l-glutamine, and 50 U/ml penicillin–50 µg/ml streptomycin. Unless otherwise stated, all tissue culture media was obtained from Life Technologies. The NaV1.9 band was normalized to the transferrin band for protein loading. Western blot analysis was performed by incubation for 1 h at RT with anti-FLAG diluted 1:15,000 for NaV1.9 detection or anti-transferrin (GIBCO/Invitrogen) diluted 1:5,000 as a gel loading control. The secondary antibody (goat anti–mouse, BioTechnology, Inc.) was horseradish peroxidase conjugated and detection or anti-transferrin (GIBCO/Invitrogen) diluted 1:5,000 as a gel loading control. The secondary antibody (goat anti–mouse, same for both primary antibodies, diluted 1:10,000; Santa Cruz Biotechnology, Inc.) was horseradish peroxidase conjugated and incubated at RT for 1 h. Densitometry of protein bands was used to quantify protein levels (ImageJ software; National Institutes of Health). The NaV1.9 band was normalized to the transferrin band to normalize for protein loading.

Plasmids and cell transfection
A full-length, human NaV1.9 (GenBank accession no. NM_014139) cDNA including a C-terminal triple FLAG epitope (SRDYKDHDGYKDHDIDYKDDDDKSR) was synthesized by DNA2.0 Inc. without codon optimization and cloned into a low copy number plasmid (pJ251). The FLAG epitope does not prevent plasma membrane trafficking of other neuronal Na channel proteins expressed heterologously or in vivo (Kearney et al., 2001; Kahlig et al., 2008). Stable expression of NaV1.9-3xFLAG in ND7/23 cells was achieved using the piggyBac transposon system as previously described (Wilson et al., 2007; Kahlig et al., 2010) using a transposon vector containing a puromycin resistance gene (pB-NaV1.9-3xFLAG-PuroR). ND7/23 cells were transfected with pCMV-hyPBase encoding a hyperactive version of the piggyBac transposase (Yusa et al., 2011) with pB-NaV1.9-3xFLAG-PuroR using FUGENE-6 (Roche). Stable NaV1.9-expressing cells were selected with 5 µg/ml puromycin (GIBCO/Invitrogen), and individual cell colonies were isolated using cloning discs and expanded.

NaV1.9 protein expression
Cells were incubated at 37°C or 28°C with 5% CO₂ for ~24 h before protein isolation. NaV1.9 was detected using a monoclonal anti-FLAG antibody (Sigma-Aldrich). NaV1.9-expressing cells (60-mm dishes) were washed twice with Dulbecco’s PBS supplemented with CaCl₂ and MgCl₂ and then incubated with 1.5 mg/ml membrane-impermeant sulfo-NHS-biotin reagent (Thermo Fisher Scientific) for 1 h on ice. The cells were lysed with 500 µl RIPA buffer (150 mM NaCl, 50 mM Tris-Base, pH 7.5, 1% IGEPAL, 0.5% Na deoxycholate, and 0.1% SDS) supplemented with a Complete mini protease inhibitor tablet (Roche) and centrifuged at 14,000 g for 30 min at 4°C, and the supernatant fraction was collected and retained as the total lysate protein fraction. Protein lysates were quantified using the Bradford reagent (Bio-Rad Laboratories), and equal amounts of proteins were used in the immunoprecipitation experiments. The lysates were incubated with 50 µl ImmunoPure Immobilized Streptavidin beads (Thermo Fisher Scientific) overnight at 4°C. The samples were centrifuged for 10 s at 10,000 g and then washed three times with RIPA buffer at 4°C. The biotinylated proteins were eluted with 57 µl Laemmli sample buffer (Bio-Rad Laboratories) supplemented with 5% 2-mercaptoanol (β-ME) for 30 min at room temperature (RT). Western blot analysis was performed by incubation for 1 h at RT with anti-FLAG diluted 1:15,000 for NaV1.9 detection or anti-transferrin (GIBCO/Invitrogen) diluted 1:5,000 as a gel loading control. The secondary antibody (goat anti–mouse, same for both primary antibodies, diluted 1:10,000; Santa Cruz Biotechnology, Inc.) was horseradish peroxidase conjugated and incubated at RT for 1 h. Densitometry of protein bands was used to quantify protein levels (ImageJ software; National Institutes of Health). The NaV1.9 band was normalized to the transferrin band to normalize for protein loading.

Electrophysiology
Cells were incubated at 28°C with 5% CO₂ for ~24 h before their use in electrophysiology experiments. Cells were dissociated by trituration, resuspended in supplemented DMEM medium (without puromycin), plated on glass coverslips, and allowed to recover for ~2 h before electrophysiological experiments. Cells were first maintained for 1 h at 37°C in 5% CO₂ and then at 28°C with 5% CO₂ until used in the experiments. For single-channel experiments, the glass coverslips were pretreated with CELL-TAK cell and tissue adhesive (Collaborative Biomedical Products).

Na currents were recorded at RT in the whole cell or excised, outside-out patch configuration of the patch-clamp technique (Hamill et al., 1981) using an Axopatch 200B series amplifier (Molecular Devices). Whole-cell currents were acquired at 20 kHz and filtered at 5 kHz. Pulse generation and data collection were performed with Clampex 9.2 (Molecular Devices). Bath solution contained (in mM) 145 NaCl, 4 KCl, 1.8 CaCl₂, 1 MgCl₂, and 10 HEPES (N-(2-hydroxyethyl)piperazine-N'-2-ethanesulphonic acid), pH 7.35, 310 mOsm/kg. 150 nM TTX was present in the bath to block endogenous Na currents. The

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composition of the pipette solution was (in mM) 10 NaF, 110 CsF, 20 CsCl, 2 EGTA (ethyleneglycol-bis-(β-aminoethylthylether), and 10 HEPES, pH 7.35, 310 mOsm/kg. Osmolarity and pH values were adjusted with sucrose and NaOH, respectively. Whole-cell patch pipettes were pulled from thin-wall borosilicate glass, and single-channel patch pipettes were pulled from standard-wall borosilicate glass (Warner Instruments, LLC) with a multistage P-97 Flaming-Brown micropipette puller (Sutter Instrument) and fire-polished with a Micro Forge MF 830 (Narashige). Pipette resistance was ~2 MΩ for whole-cell recording and 4–6 MΩ for outside-out patch recording. Single-channel recording pipettes were coated with Sylgard 184 (Dow Corning). 2% agar bridges containing bath solution served as reference electrodes. Cells were allowed to equilibrate for 10 min in bath solution before obtaining seals. Peak currents were measured using 50-ms pulses to between −100 and 40 mV every 5 s from a holding potential of −120 mV. The peak current was normalized for cell capacitance and plotted against voltage to generate peak current density–voltage relationships. For recording of NaV1.9 currents in low Na+ bath solution ([Na+]o = 20 mM), NaCl was replaced with equimolar N-methyl-d-glucamine-chloride (NMDG-Cl). The liquid junction potential generated upon reducing extracellular Na+ was calculated using the Junction Potential Calculator in Clampex 9.2. The calculated junction potential was 4.5 mV. Whole-cell conductance was calculated from the peak current amplitude using the formula $G_{Na} = I_{peak} / (V - E_{rev})$, where $E_{rev}$ is the estimated Na+ reversal potential, and then normalized to the maximal conductance recorded between Na+ reversal potential, and then normalized to the maximal conductance recorded between −80 and 20 mV. The normalized G–V curves were fit with the Boltzmann function, $G = G_{max} \times \frac{(1 + \exp(V - V_{1/2})/k)}{(1 + \exp(V - V_{1/2})/k)}$, to determine the voltage for half-maximal channel activation ($V_{1/2}$) and slope factor ($k$). The voltage dependence of channel availability was assessed after a 300-ms prepulse to various potentials followed by 50-ms pulse to −40 mV, the voltage at which peak NaV1.9 currents were measured. The normalized current was plotted against the voltage, and the steady-state channel availability curves were fit with Boltzmann functions ($I/Imax = 1/(1 + \exp(V - V_{1/2})/k)$) to determine the voltage for half-maximal channel inactivation ($V_{1/2}$) and slope factor ($k$). Recovery from fast inactivation was examined as fractional recovery against the recovery period and fitted to a five-parameter double exponential function, $I/Imax = A_0 \times \left[ 1 - \exp(-\tau_{1}/C) \right] + A_1 \times \left[ 1 - \exp(-\tau_{2}/C) \right] + A_2 \times \left[ 1 - \exp(-\tau_{3}/C) \right]$, where $\tau_1$ and $\tau_2$ denote time constants (fast and slow components, respectively) and $A_0$, $A_1$, and $A_2$ represent the fast and slow fractional amplitudes. Single-channel behavior was examined during a −40-mV pulse from a holding potential of −120 mV in the presence of 150 nM external TTX. Single-channel currents were filtered at 2 kHz and acquired at 20 kHz. Each test pulse was followed by a 10-s recovery period at −120 mV. Linear leak and capacitive transients were subtracted using the mean of blank traces, and no corrections were made for missed events. The number of channels in each patch was determined as the maximum number of simultaneously open channels observed in all sweeps at a given voltage step. Only membrane patches without channel activity at the holding potential (~120 mV) or outward openings at the test potential (~40 mV) were analyzed, and this excluded contaminating currents from the other possible charge carriers present in the solutions (i.e., Ca2+, F−, and Cl−).

**Data analysis**

Data were analyzed and plotted using a combination of Clampfit 9.2 (Molecular Devices), SigmaPlot 10 (Systat Software, Inc.) and Origin 7.0 (OriginLab). Statistical analyses were performed using SigmaStat 2.03 (Systat Software, Inc.), and statistical significance was determined using unpaired Student’s t test. Data are represented as means ± standard error of the mean. The number of cells/patches used for each experimental condition is given in the figure legends.

**RESULTS**

**Stable expression of human NaV1.9**

We generated a stable cell line expressing full-length, FLAG-tagged human NaV1.9 cDNA (NaV1.9-3xFLAG) in ND7/23 cells using a previously described transposon system (Wilson et al., 2007; Kahlig et al., 2010). We selected ND7/23 cells because of their chimeric derivation from rat DRG and mouse neuroblastoma cells and because our attempts to generate stably transfected HEK-293 and Chinese hamster ovary cells were unsatisfactory.

Initial electrophysiological analysis of NaV1.9-expressing cells (performed in the presence of 150 nM TTX to block an endogenous TTX-sensitive NaV channel) demonstrated small TTX-resistant currents (Fig. 1A, top). Based on prior reports that low-temperature incubation of heterologous cells increased the functional expression of ion channels (e.g., CFTR and hERG; Denning et al., 1992; Zhou et al., 1999), we tested this strategy with NaV1.9. Overnight incubation of NaV1.9-ND7/23 cells at 28°C increased the current magnitude considerably (Fig. 1A, A and B), whereas incubation of nontransfected ND7/23 cells at 28°C did not generate TTX-resistant currents. The NaV1.9-dependent currents were markedly attenuated when the extracellular Na+ concentration was reduced, consistent with an Na-selective channel (peak current: 145 mM external [Na+], 37.4 ± 13.2 pA/pF; 20 mM external [Na+], 6.5 ± 1.9 pA/pF; n = 5). Further, application of 30 μM TTX blocked peak current recorded at −40 mV by a mere 12.8 ± 2.9% (n = 4), consistent with a TTX-resistant channel.

Biochemical analysis of total and cell surface protein isolated from NaV1.9 stable cells cultured at 37°C and 28°C showed a prominent immunoreactive band (~205 kD; Fig. 1C) approximately the size predicted for native NaV1.9. Previous analysis of NaV1.9 protein from adult rat DRG neurons indicated that the channel is lightly glycosylated (Tyrrell et al., 2001), and this is consistent with our observation. Densitometry analysis of NaV1.9 immunoblots (Fig. 1C) demonstrated that total and cell surface NaV1.9 protein expression were increased approximately twofold after overnight incubation of the cells at 28°C, but the molecular mass did not change (Fig. 1D). The mean ratio of surface to total protein trended higher for cells incubated at 28°C (37°C = 0.8 ± 0.3; 28°C = 1.1 ± 0.3; n = 3), but the difference was not statistically significant.

**Biophysical properties of human NaV1.9**

After establishing stable heterologous expression of NaV1.9, we proceeded with determining more detailed functional properties of this unusual Na channel. As illustrated in Fig. 1A, the kinetics of channel gating, especially inactivation, are markedly slow and have substantial voltage dependence. Furthermore, inactivation
measured recovery from inactivation after a 300-ms depolarization pulse to ~40 mV. The recovery rate exhibited two exponential components with a predominant fast and a smaller slow component. Quantified biophysical parameters for the voltage dependence of activation, steady-state inactivation, and recovery from fast inactivation are provided in Table 1.

**Na\textsubscript{v}1.9 activation by GTP\textsubscript{S}\textsubscript{Y}**

Inflammatory mediators acting through G-protein-coupled receptors are known to potentiate Na\textsubscript{v}1.9 channel activity and increase TTX-resistant persistent Na current in sensory neurons (Baker et al., 2003; Maingret et al., 2008b; Ostman et al., 2008; Ritter et al., 2009). These effects can be mimicked by intracellular application of GTP\textsubscript{S}\textsubscript{Y} (a hydrolysis-resistant GTP analogue), and we tested whether this phenomenon could be replicated in Na\textsubscript{v}1.9-expressing ND7/23 cells. Fig. 3A illustrates the mean whole-cell currents recorded from Na\textsubscript{v}1.9-expressing cells in the absence and presence of 200 µM of intracellular GTP\textsubscript{S}\textsubscript{Y}. Application of intracellular GTP\textsubscript{S}\textsubscript{Y} had a concentration-dependent effect on Na\textsubscript{v}1.9 current density (Fig. 3B and C). Addition of 20 µM GTP\textsubscript{S}\textsubscript{Y} had no effect, whereas 63 µM increased the current density by 40% and 200 µM approximately doubled Na\textsubscript{v}1.9 current density (Fig. 3B and C). All subsequent experiments were performed with 200 µM GTP\textsubscript{S}\textsubscript{Y}. The doubling of Na\textsubscript{v}1.9 current density by GTP\textsubscript{S}\textsubscript{Y} was evident 2 min after rupture of the membrane patch and intracellular dialysis with the pipette solution (Fig. 3D). The increase in current density was accompanied by a depolarizing shift in the voltage dependence of inactivation without any effect on the

![Figure 1](image-url)

**Figure 1.** Incubation at 28°C increases Na\textsubscript{v}1.9 cell surface expression and current magnitude. (A) Averaged whole-cell currents recorded from cells incubated at 37°C (n = 8) or 28°C (n = 9). (B) Current-voltage relationships measured from Na\textsubscript{v}1.9-expressing cells at 37°C or 28°C. (C) Total and cell surface Na\textsubscript{v}1.9 protein isolated from cells grown at 37°C or 28°C. (D) Densitometric analysis of total and cell surface Na\textsubscript{v}1.9 bands obtained from three different protein isolations. Data are represented as means ± standard error of the mean.
dependence of activation and increased the time constant corresponding to the slow component of recovery from fast inactivation (Table 1). Whole-cell activation kinetics were not affected by intracellular GTP\textsubscript{S}; both activation time constants (Fig. 4 A) and time to peak (Fig. 4 B) were similar to control values. However, GTP\textsubscript{S} did affect inactivation kinetics at potentials voltage dependence of activation or the time course of recovery from inactivation (Fig. 3, E and F; and Table 1). Importantly, application of intracellular GTP\textsubscript{S} did not evoke expression of TTX-resistant currents in nontransfected ND7/23 cells.

More detailed analyses revealed that intracellular GTP\textsubscript{S} slightly decreased the slope factor for voltage dependence of activation and increased the time constant corresponding to the slow component of recovery from fast inactivation (Table 1). Whole-cell activation kinetics were not affected by intracellular GTP\textsubscript{S}; both activation time constants (Fig. 4 A) and time to peak (Fig. 4 B) were similar to control values. However, GTP\textsubscript{S} did affect inactivation kinetics at potentials

<table>
<thead>
<tr>
<th>Condition</th>
<th>Peak current density (−40 mV)</th>
<th>Voltage dependence of activation</th>
<th>Steady-state inactivation</th>
<th>Recovery from inactivation</th>
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<tr>
<td></td>
<td>pA/pF</td>
<td>V\textsubscript{1/2}</td>
<td>k</td>
<td>n</td>
</tr>
<tr>
<td>Control</td>
<td>−23.2 ± 3.4</td>
<td>16</td>
<td>−56.9 ± 0.6</td>
<td>6.7 ± 0.3</td>
</tr>
<tr>
<td>GTP\textsubscript{S}</td>
<td>−38.9 ± 5.1 (^a)</td>
<td>20</td>
<td>−54.3 ± 1.1</td>
<td>5.8 ± 0.3 (^b)</td>
</tr>
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\(^{a}\)P = 0.025 (GTP\textsubscript{S} vs. control using Student’s t test).

\(^{b}\)P = 0.015 (GTP\textsubscript{S} vs. control using Student’s t test).

\(^{c}\)P = 0.021 (GTP\textsubscript{S} vs. control using Student’s t test).

\(^{d}\)P = 0.011 (GTP\textsubscript{S} vs. control using Student’s t test).

Steady-state inactivation was determined using a 300-ms prepulse to various potentials followed by a 50-ms pulse to −40 mV. Recovery from inactivation was determined after a 300-ms prepulse to −40 mV.

\(^{a}\)P = 0.025 (GTP\textsubscript{S} vs. control using Student’s t test).

\(^{b}\)P = 0.015 (GTP\textsubscript{S} vs. control using Student’s t test).

\(^{c}\)P = 0.021 (GTP\textsubscript{S} vs. control using Student’s t test).

\(^{d}\)P = 0.011 (GTP\textsubscript{S} vs. control using Student’s t test).

Table 1: Biophysical properties of Na\textsubscript{v}1.9 in the absence (control) or presence of intracellular GTP\textsubscript{S}
positive to −50 mV. Fig. 4 C shows whole-cell current inactivation of Na_{1.9} in the presence of intracellular GTPγS. Specifically, the inactivation time course determined between −40 mV and 10 mV exhibited a biexponential decay (Fig. 4 D) with voltage-dependent time constants and fractional amplitudes. At −40 mV, both the slow and fast components were equally proportioned, whereas at more positive potentials the fast component fraction predominated (≈65% at −20 mV and ≈80% at 10 mV, Fig. 4 E). These data indicated that GTPγS evoked changes in Na_{1.9} gating, but these findings from whole-cell recording experiments did not provide a clear explanation for increased current amplitude.

Single-channel mechanism of Na_{1.9} potentiation by GTPγS

To further elucidate biophysical mechanisms that explain GTPγS potentiation of Na_{1.9} current, we performed single-channel analysis on Na_{1.9}-ND7/23 cells in the absence or presence of intracellular GTPγS. These experiments were performed in the presence of 150 nM external TTX to block endogenous Na_{1.9} channels.

Fig. 5 illustrates representative single-channel traces recorded at −40 mV from excised outside-out membrane patches dialyzed with either control intracellular solution or with solution containing 200 µM GTPγS. Single-channel amplitudes were similar for both conditions (control, 0.69 ± 0.05 pA; +GTPγS, 0.72 ± 0.04 pA; Fig. 5 C). This observation excluded a fundamental change in ion permeation as the cause for increased whole-cell current amplitude evoked by GTPγS. In contrast, exposure of Na_{1.9} to intracellular GTPγS was associated with an ≈2.7-fold larger nPo (control, 0.23 ± 0.07, n = 4; +GTPγS, 0.63 ± 0.05, n = 4; P = 0.004; Fig. 5 D) and a significantly greater mean open time deduced from open time histograms (Fig. 5 E). By dividing the nPo value calculated for each outside-out patch by the maximum number of channels observed in the same patch, we calculated an ≈3.3-fold larger Po for channels in the presence of GTPγS (Po: control = 0.06 ± 0.01 [4.0 ± 1.3 channels/patch]; +GTPγS = 0.19 ± 0.02 [3.2 ± 0.2 channels/patch]). These findings indicated that changes in gating were primarily responsible for GTPγS potentiation of Na_{1.9} current.
Our success in achieving stable heterologous expression of NaV1.9 can be attributed to three factors. First, we engineered NaV1.9 stable cells using a highly efficient transposon system capable of integrating transgenes into multiple loci within a host cell genome (Kahlig et al., 2010). This approach may also bias integration events to regions of active chromatin increasing the likelihood of transgene expression (Wilson et al., 2007). Second, use of ND7/23 cells may provide intracellular factors expressed in neurons that are absent in other commonly used heterologous cells (e.g., HEK-293). For example, contactin has been demonstrated to bind to NaV1.9 and increase surface expression and current density of NaV1.9 channels in DRG neurons (Liu et al., 2001; Rush et al., 2005). Finally, we used low-temperature (28°C) incubation of cells to promote higher levels of functional NaV1.9 expression. The efficacy of 28°C incubation to boost channel activity may reflect a certain degree of impaired NaV1.9 trafficking in heterologous cells. However, our biochemical data did not demonstrate a clear change in the ratio of cell surface to total cellular protein consistent with

**DISCUSSION**

In this study, we sought to determine the functional properties of recombinant human NaV1.9 and the mechanism by which G-protein signaling increases NaV1.9 activity. Because this NaV channel has been implicated in nociception, our findings have implications for understanding the physiology of pain and could provide the basis for new analgesic strategies.

**Heterologous functional expression of human NaV1.9**

Our experiments were enabled by successful and robust stable expression of human NaV1.9 in heterologous cells. Previously, heterologous expression of NaV1.9 has been difficult (Dib-Hajj et al., 2002), leading to speculation this is a not a voltage-gated channel, but rather a ligand-gated channel (Blum et al., 2002). However, the absence of TTX-resistant persistent Na current in mouse Scn11A-null neurons and its restoration by transfection of human NaV1.9 in these cells (Ostman et al., 2008) eliminated doubt regarding the true nature of this protein.

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**Figure 4.** Effects of intracellular GTP\(\gamma\)S on NaV1.9 kinetics. (A) Time constants for activation in the presence of intracellular GTP\(\gamma\)S \((n = 13–14)\). (B) Time to peak current in the presence of intracellular GTP\(\gamma\)S \((n = 21)\). Dotted lines illustrate data for control solution (from Fig. 2). (C) Averaged whole-cell NaV1.9 currents recorded during 300-ms pulses to \(-80, -60, -40, -20\), and \(0\) mV. Dotted line indicates the zero current level \((n = 10)\). (D) Inactivation time constants (fast, blue; slow, red) recorded in the presence of intracellular GTP\(\gamma\)S \((n = 8–14)\). Inactivation time constants for control pipette solution (open circles; \(n = 13–17\)) are also shown (from Fig. 2 D). (E) Fractional amplitudes for inactivation time constants determined in the presence of intracellular GTP\(\gamma\)S for \(-40\) to \(10\) mV \((n = 8–14)\). Data are represented as means ± standard error of the mean.
redistribution of channels to the plasma membrane. Alternative explanations for the effects of low temperature include more efficient translation of NaV1.9 or slower protein turnover. These various maneuvers appear to overcome the natural tendency for NaV1.9 to express small Na currents, an intrinsic property of the channel which may protect cells from intracellular Na+ overload or excessive membrane depolarization.

Biophysical properties of human NaV1.9

Using stably expressing heterologous cells, we determined the biophysical properties of human NaV1.9 using both whole-cell and single-channel recording strategies. Among the more characteristic features of NaV1.9 is its unusually slow gating, especially inactivation. In single-channel recordings, long openings, and tendency for re-openings account for the sluggish nature of inactivation. This characteristic of NaV1.9 inactivation stems in part from the large overlap between the voltage dependence of activation and inactivation with an expanded range of voltages in which channels are active but not subject to strong inactivation tendencies (“window” current). Furthermore, the voltage midpoint (V1/2) for activation (−56 mV) is similar to that for inactivation (−52 mV). In contrast, fast-gating NaV channels have inactivation V1/2 values that are typically more negative than activation. The effect of this relationship is illustrated at −80 mV (Fig. 2 C), a voltage at which NaV1.9 is activated but exhibits very little inactivation. Our findings are in agreement with previous experiments associating this channel with subthreshold Na current observed in sensory neurons. Some species-related differences in the reported voltage dependence of activation and inactivation for NaV1.9 are evident when comparing our data with recordings from rats and mice (Cummins et al., 1999; Coste et al., 2004; Rush and Waxman, 2004), but the general trends are similar. However, the structural basis for the unusual gating behavior of NaV1.9 is unclear.

Figure 5. Effect of GTPγS on NaV1.9 single-channel activity. Single NaV1.9 channel openings were elicited by 300-ms pulses to −40 mV from a holding potential of −120 mV with 150 nM TTX present in the bath solution. (A) Single-channel traces from excised, outside-out patches dialyzed with control intracellular solution. Dotted lines indicate the zero current level, and individual single-channel amplitudes are marked by horizontal lines. The illustrated patch recordings contain at least six channels. (B) Single-channel traces from excised, outside-out patches dialyzed with pipette solution containing 200 µM GTPγS. The illustrated patch recordings contain at least four channels. (C) Single-channel amplitudes measured in the absence (control, n = 4) or presence (+GTPγS, n = 4) of 200 µM GTPγS. (D) Values for nPo calculated from single-channel recordings from control and +GTPγS patches. Intracellular GTPγS increased nPo ~2.5 (*, P = 0.004). (E) Open time histograms plotted for single-channel recordings performed in the absence or presence of GTPγS. The mean open times calculated for control solution were τ1 = 0.09 ± 0.01 ms (A1 = 95.7 ± 1.8%) and τ2 = 0.47 ± 0.07 ms (A2 = 4.3 ± 1.8%; n = 4). In the presence of GTPγS, mean open times were τ1 = 0.24 ± 0.05 ms (A1 = 91.5 ± 2.4%) and τ2 = 1.55 ± 0.35 ms (A2 = 8.5 ± 2.4%; n = 4). The differences in τ1 and τ2 values between control and +GTPγS were significant, P = 0.026 and P = 0.023, respectively. Data are represented as means ± standard error of the mean.
Mechanisms of GTPγS potentiation of human NaV1.9

Previous studies have demonstrated that G-protein activation increases TTX-resistant, persistent Na current in sensory neurons and that this phenomenon requires NaV1.9 (Baker et al., 2003; Rush and Waxman, 2004; Ostman et al., 2008). In this study, we demonstrated that heterologously expressed human NaV1.9 can also be potentiated by intracellular GTPγS, a nonselective activator of G-protein signaling. We observed that exposure to GTPγS caused an approximate twofold greater current density accompanied by an ~10-mV depolarizing shift in the voltage dependence of inactivation but without a change in activation voltage dependence. The unchanged voltage dependence of activation in combination with a more positive voltage dependence of inactivation yields a larger window current after GTPγS exposure. Additionally, GTPγS slowed inactivation by introducing a slower component readily discernable in a physiological voltage range. In contrast, study of murine NaV1.9 demonstrated that NaV1.9 potentiation by GTPγS exposure closely approximates a major change in permeation properties as the source for increased current density evoked by GTPγS. However, intracellular GTPγS did cause a significantly greater mean open time, which could account for the slower time course of inactivation and thus larger persistent current evoked by G-protein activation. We also examined whether channel open probability was affected. We determined that nPo was ~2.7 times greater in the presence than absence of intracellular GTPγS. Further analysis revealed that a greater Po associated with GTPγS exposure closely approximates the fold increase in nPo and whole-cell current density. Together, these results indicated that G-protein activation potentiates human NaV1.9 by increasing channel open probability and lengthening mean open time, causing the larger peak and persistent current after exposure to inflammatory mediators.

In summary, we have developed a stable cell line expressing human NaV1.9 and overcome previous challenges to investigating this channel in heterologous cells.

We analyzed the biophysical properties of human NaV1.9 and determined the mechanism by which GTPγS potentiates channel activity. Our results demonstrate a cellular platform that will be useful for the discovery of NaV1.9 blockers or modulators and to understand how inflammatory agents regulate NaV1.9 current. Furthermore, our findings may contribute to the development of new treatment strategies for chronic and inflammatory pain.

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