Modulation of human Nav1.7 channel gating by synthetic α-scorpion toxin OD1 and its analogs

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ABSTRACT
Nine different voltage-gated sodium channel isoforms are responsible for inducing and propagating action potentials in the mammalian nervous system. The Nav1.7 channel isoform plays an important role in conducting nociceptive signals. Specific mutations of this isoform may impair gating behavior of the channel resulting in several pain syndromes. In addition to channel mutations, similar or opposite changes in gating may be produced by spider and scorpion toxins binding to different parts of the voltage-gated sodium channel. In the present study, we analyzed the effects of the α-scorpion toxin OD1 and 2 synthetic toxin analogs on the gating properties of the Nav1.7 sodium channel. All toxins potently inhibited channel inactivation, however, both toxin analogs showed substantially increased potency by more than one order of magnitude when compared with that of wild-type OD1. The decay phase of the whole-cell Na⁺ current was substantially slower in the presence of toxins than in their absence. Single-channel recordings in the presence of the toxins revealed that Na⁺ current inactivation slowed due to prolonged flickering of the channel between open and closed states. Our findings support the voltage-sensor trapping model of α-scorpion toxin action, in which the toxin prevents a conformational change in the domain IV voltage sensor that normally leads to fast channel inactivation.

KEYWORDS
gating; Nav1.7; patch clamp; scorpion toxin; voltage-gated sodium channel

Introduction
The Na+,1.7 voltage-gated sodium channel (VGSC) is encoded by SCN9A; expressed in sensory and sympathetic neurons, Schwann and endocrine cells; and plays a key role in generating and propagating action potentials in the central and peripheral nervous systems. Na+,1.7 channel mutations may result in gain- or loss-of-function, creating pathological conditions such as over-sensitivity or loss of sensitivity to pain, respectively. For example, the S241T and A1746G mutations, which have been linked to inherited or primary erythromelalgia, result in a shift in voltage-dependence of activation to more negative potentials that makes cells hyper-excitiable.

Certain toxins from animal venoms can cause similar effects to those caused by mutations. For instance, some VGSC-targeting spider and scorpion β-toxins frequently bind to receptor site 4 (on domain II), where they can shift voltage-dependence of activation to more negative potentials (‘activators’) or more positive potentials (‘inhibitors’). Toxins targeting site 3 (on domain IV), such as spider toxins, δ-conotoxins and α-scorpion toxins, typically slow inactivation, which prolongs the decay phase of the whole-cell Na⁺ current. Interestingly, some toxins appear to target sites 3 and 4 simultaneously, affecting fast inactivation and steady-state activation. A classical example is spider toxin ProTx-II, which has been shown to interact with DII and DIV on Na+,1.7, although the affinities are substantially different. Another example is the recently characterized scorpion α-toxin OD1, which potently inhibits mammalian Na+,1.4, Na+,1.6 and Na+,1.7 inactivation and results in a hyperpolarizing shift in the steady-state voltage-dependence of Na+,1.4 and Na+,1.6 activation.

Using a high-throughput fluorescence-based assay, we recently identified OD1 analogs with altered Na+,1.4,
Nav1.6 and Nav1.7 selectivity, and significantly improved potency at Nav1.7. This assay relies on membrane potential-sensitive dyes to monitor Na₉ activation in the presence of sub-threshold concentrations of the site 2-targeting alkaloid veratridine, and facilitates rapid, qualitative and quantitative analysis of VGSC modulators. However, the lack of control over the membrane potential in this assay precludes any detailed, mechanistic VGSC studies at the molecular level.

Here, we report a detailed analysis of how mutations of OD1 affect the toxin’s interaction with the Nav1.7 VGSC.

**Results**

**Concentration-dependent effect of OD1 and analogs on sodium current inactivation**

Step depolarization of CHO cells stably expressing human Nav1.7 channels from −100 mV to 0 mV in the absence of toxins elicited the whole-cell inward Na⁺ current with fast onset, followed by an exponential decay with time constants τ₁ = 0.77 ± 0.03 ms and τ₂ = 6.0 ± 0.4 ms (n = 6). Applying OD1 and its analogs remarkably prolonged the current decay phase in a concentration-dependent manner (Fig. 1A). This effect was quantified by the ratio of the current measured at an interval of 1.5 ms from the current peak to the peak value (Fig. 1B).

The potency of the OD1 analogs was substantially higher than that of wild-type OD1, exceeding it approximately 30 times for OD1-KPH and approximately 16 times for OD1-KPH-E55A (Fig. 1B). Effects of the toxins on inactivation kinetics were completely eliminated by toxin washout or upon applying strong depolarizing pre-pulses (up to +190 mV) in the presence of OD1-KPH (Fig. 2). The degree of current recovery depended on the amplitude of depolarization, and was estimated by fitting the experimental data with a logistic equation (Fig. 1B).

**Figure 1.** Wild-type OD1 and its analogs KPH and KPH-E55 slow the decay phase of whole-cell Na⁺ current in a concentration-dependent manner. (A) Three sets of normalized current traces represent responses to step depolarization from a holding potential of −100 mV to 0 mV. Each set consists of a trace in the absence of the toxin (upper fast declining current), at approximately EC₅₀ concentration (middle trace) and saturating concentration. The concentrations were as follows: OD1: 0, 10 nM, 300 nM; KPH: 0, 300 pM, 10 nM; and KPH-E55A: 0, 300 pM, 10 nM. (B) Concentration-dependence of change in inactivation rate of whole-cell Na⁺ currents. The ratio of whole-cell value taken at 1.5 ms after the peak occurrence to the peak current was taken as an estimate of inactivation rate. Experimental points were fitted with a logistic equation with the following parameters: OD1: EC₅₀ = 12 ± 3 nM, n = 0.7 ± 0.2; KPH: EC₅₀ = 0.4 ± 0.1 nM, n = 1.0 ± 0.3; and KPH-E55A: EC₅₀ = 0.77 ± 0.08 nM, n = 1.0 ± 0.1.
data to a logistic function yielding a midpoint value of $+127 \pm 1$ mV ($n = 4$) (Fig. 2B). Pre-pulses to potentials more positive than $+190$ mV were detrimental to the cell membrane and were not examined.

**Effects of OD1 and analogs on Na$_{\text{v}}$.1.7 channel gating**

We analyzed the effects of OD1 and its analogs on Na$_{\text{v}}$.1.7 voltage-dependence of activation and inactivation. Current–voltage (I–V) relationships in the absence and presence of toxins are shown in Figure 3A. Half-activation voltage ($V_{0.5}$) for Na$_{\text{v}}$.1.7 was $-29.3 \pm 0.3$ mV ($n = 14$) in the absence of toxin, and leftward shifted to more negative potentials in the presence of 10 nM wild-type OD1 ($V_{0.5} = -35.8 \pm 0.3$ mV, $n = 7$), 500 pM OD1-KPH ($V_{0.5} = -37.9 \pm 0.2$ mV, $n = 5$) or 1 nM OD1-KPH-E55A ($V_{0.5} = -39.7 \pm 0.2$ mV, $n = 7$), respectively. The minor, but statistically significant, shift caused by wild-type OD1 is in agreement with earlier observations by Maertens et al.,$^{15}$ who observed a $-3$ mV shift.

Along with I–V changes, the toxins caused similar shifts in the voltage-dependence of activation (Fig. 3B, Table 1). The resulting $V_{0.5}$ shifts are minor and, at least for wild-type OD1, substantially smaller than the negative shifts observed on Na$_{\text{v}}$.1.4 ($-13$ mV) and Na$_{\text{v}}$.1.6 ($-20$ mV).$^{16}$ In contrast with their effect on steady-state activation, neither

|                     | $G/G_{\text{max}}$ | $I/I_{\text{max}}$ |
|---------------------|-------------------|------------------|
|                     | $V_{0.5}$ (mV)    | $n$   | $p$   | $V_{0.5}$ (mV) | $n$   | $p$   |
| Control             | $-29.1 \pm 0.2$   | 14    |  —   | $-72.2 \pm 0.4$ | 14    |  —   |
| OD1 (10 nM)         | $-35.2 \pm 0.3$   | 7     | 0.004| $-66.7 \pm 0.4$ | 5     | 0.1   |
| KPH (0.5 nM)        | $-35.3 \pm 0.1$   | 5     | 0.005| $-69.9 \pm 0.2$ | 5     | 0.3   |
| KPH-E55A (1 nM)     | $-39.1 \pm 0.3$   | 7     | 0.0001| $-69.5 \pm 0.4$ | 7     | 0.3   |

$P$ values calculated with Student t test against control.

Figure 2. Recovery of Na$^+$ current from toxin action by strong depolarization and washout. (A) Whole-cell Na$^+$ current elicited by membrane depolarization to 0 mV in the presence of 500 pM KPH (black traces). Strong depolarizing pre-pulse to $+190$ mV restores inactivation rate (red trace on top inset). Na$^+$ currents showed fast and complete recovery from toxin action within 5 min of washout (red trace on the bottom inset). Similar results were obtained for OD1 and KPH-E55A. (B) Voltage-dependence of toxin displacement by depolarizing pre-pulse in the presence of 500 pM KPH ($n = 4$).
OD1-KPH nor OD1-KPH-E55A have a statistically significant effect on the Na$_v$1.7 voltage-dependence of fast inactivation (Fig. 3B).

**Recovery from fast inactivation**

OD1 and its analogs KPH and KPH-E55A enhance Na$_v$1.7 recovery from fast inactivation (Fig. 4). The recovery rate was estimated from the ratio of peak current evoked by a 40 ms test pulse to 0 mV, and peak current recorded during the preceding 40 ms pre-pulse to 0 mV, as a function of time interval between these 2 pulses. Data obtained in the absence of the toxin could be successfully fit by a single exponential function with a time constant of $\tau = 15 \pm 2$ ms ($n = 14$).

In contrast, applying all toxins resulted in pronounced biphasic recovery from inactivation kinetics, with both kinetically distinct phases having approximately equal amplitudes. Experimental points were fit by a double exponential function. Parameters of the fits are given in Table 2.

**Single Na$^+$ channel recordings**

It is well established that the observed whole-cell current waveform may be realized by substantially different molecular mechanisms of gating. To get a deeper insight into these mechanisms, we investigated the action of OD1 at the microscopic level using single-channel patch clamp recording. Representative examples of single-channel Na$_v$1.7 currents from outside-out patches are shown in Figure 5A. Applying wild-type OD1 (100 nM) did not affect the current amplitude, which was $1.4 \pm 0.2$ pA and $1.36 \pm 0.09$ pA ($n = 3$) in the absence and presence of OD1, respectively. Surprisingly, the mean open time of the channels also was not substantially changed by the toxin (Fig. 5B). In contrast, Na channels exhibited flickering behavior in the presence of the toxin, which

**Table 2.** Amplitudes ($A$) and time constants ($\tau$) of recovery from fast inactivation state.

|       | $A_i$ (pA) | $\tau_i$ (ms) | $A_f$ (pA) | $\tau_f$ (ms) | n |
|-------|------------|---------------|------------|---------------|---|
| Control | 0.94 ± 0.02 | 15 ± 2 | — | — | 14 |
| OD1   | 0.48 ± 0.03 | 15 ± 2 | 0.48 ± 0.04 | 2.8 ± 0.5 | 12 |
| KPH   | 0.46 ± 0.04 | 13 ± 2 | 0.50 ± 0.05 | 2.8 ± 0.4 | 5  |
| KPH-E55A | 0.53 ± 0.05 | 15 ± 2 | 0.48 ± 0.09 | 2.5 ± 0.3 | 8  |
in some sweeps lasted for the duration of the depolarizing voltage step (Fig. 5A).

**Kinetic models**

We attempted to find an appropriate kinetic model of channel conformational states in terms of Markov’s formalism, and define the toxin’s action through the transition rates changes. As a base model (i.e. without toxin), we chose a minimal model that contained one open state, 2 closed and 2 inactivated states (Fig. 6A). Rate constants were considered stationary and not dependent on membrane potential.

To evaluate this approach, we used whole-cell experimental traces recorded in the absence of toxin under voltage-clamp conditions at a holding potential of $-100 \text{ mV}$ and a depolarizing step to $-20 \text{ mV}$. This aimed to make sure that, at the beginning of depolarization, Na channels were mostly in a ‘resting’ state $R$, and depolarizations were sufficiently strong to activate most of the channel populations. Despite a substantial reduction in the number of free parameters compared with previous studies, our base model adequately described the experimental data (Fig. 6B, upper panel).

Given our previous observations on toxin effects on steady-state channel activation (see above), we further assumed that activation pathways were not significantly affected by the toxins and placed constraints on the pathways responsible for transitioning channels from closed to open states, while leaving the others that are associated with transitioning channels to inactivated states free floating (Table 3).

The boundaries of constraints were chosen according to the rule: $|K - K_b| < \sigma$, where $K_b$ is matrix of rate constants derived from the base model, and $\sigma$ is deviation that comes from differences between experimental traces used in base model simulations. By doing so, we kept constrained rate constants close to those derived from the base model (Table 3).

To compare the different models (M1–M4) that describe the toxins’ actions, we used a logarithm error ratio (LER) used previously. Based on the LER value, model M3 best fit the experimental traces. Models that allowed transition from state C2 to state I4 in one direction and/or unconstrained transitions from state O3 to state I4 did not yield realistic values of corresponding transition rates.

**Discussion**

Similar to our previous observations, OD1 and its analogs dramatically slowed the time course of the
The apparent effect on channel inactivation is similar to that of α-scorpion toxins and has been convincingly linked to binding of the neurotoxin to site 3 on VGSCs, which comprises extracellular loops of the domain IV voltage sensor.5–7 According to the voltage-sensor trapping model, α-scorpion toxins trap the S4 transmembrane segment of the domain IV voltage sensor in the inward position and obstruct its outward movement, preventing fast inactivation.24 Single-channel recordings revealed that in the presence of toxins, channels rapidly fluctuate between open and closed states for relatively long intervals of time. On the other hand, the toxin did not affect the mean open time of the channel. This ‘flickering’ channel activity may explain the prolonged decay phase of the whole-cell Na⁺ current. The similar behavior of macroscopic currents in neuroblastoma cells affected by the α-scorpion toxin resulted from an increase in channel open time and channel reopenings.25

In addition to slowing down whole-cell current inactivation, OD1 and analogs affected the time course of the channels’ recovery from fast inactivation. In the absence of toxins, the time course of recovery followed a single exponential function, whereas an additional exponential component emerged in the presence of toxins. The time constant of the additional component was almost 5 times smaller than the recovery time constant in control conditions, and may be due to the number of channels that did not reach an inactivated state.

In terms of the Markov model, the trapping mechanism may be caused by an increase in the rate of transition from state C2 to state I4. On the other hand, the
rate constant for transition from state I4 to state O3 doubles in the presence of the toxin, making transition back to the open state more favorable. The transition rate from an open to inactivated state I4 which determines channel open time was conserved compared to that in the absence of a toxin. All of the models that we tested yielded a lower rate for the transition from the intermediate inactivated state I4 to absorbing state I5. In agreement with this mechanism, strong depolarizing pre-pulses force S4 in the DIV to move into the activated position. This might also affect the binding site for the toxin on the S3–S4 linker and abolish the toxin’s effect.8 While the importance of DIV in the channel inactivation process is well established, its involvement in the channel activation process is less clear.

The steady-state, current-voltage relationship in the presence of OD1 and its analogs showed a minor, but significant, shift to more hyperpolarized potentials. This effect is less pronounced for Na$_{1.7}$ than it is for Na$_{1.6}$ or Na$_{1.4}$, which are also affected by OD1 in a similar manner.16 It could also suggest that the DIV voltage sensors of the various isoforms are not functionally equivalent and involved in channel activation to differing degrees. However, at this stage, we cannot rule out the possibility that additional high-affinity binding (for example on DII) is responsible for the dual α/β toxin effect.

In terms of reversibility of the toxin’s action, the effect of OD1 on Na$_{1.7}$ channel inactivation differs from that of previous studies involving α-scorpion site 3 toxin Ts3 (Tityus serulatus) on Na$_{1.4}$,9,22 but with a similar effect on macroscopic current inactivation. Whereas OD1s effects on Na$_{1.7}$ are fully reversible after washout, Ts3 remains tightly bound to Na$_{1.4}$. These findings for wild-type OD1 prompted us to analyze the mechanism of action of OD1 analogs, OD1-KPH and OD1-KPH-E55A, which carry 3 or 4 non-conservative amino acid substitutions, respectively, in a structural region known as the ‘NC domain’. Both mutant toxins showed significantly improved potency in terms of their ability to inhibit fast inactivation (Fig. 1B), but from a mechanistic perspective, were otherwise indistinguishable from the wild-type toxin. This suggests that the observed increase in potency is primarily a result of an increase in binding affinity. The potency of the OD1 analogs is one order of magnitude higher than that of wild-type OD1 but there was no statistically significant difference between the V$_{0.5}$ values obtained in the presence of OD1 and its analogs.

**Materials and methods**

**Preparation of OD1 analogs**

OD1 and its triple (D9K, D10P, K11H) and quadruple mutants (D9K, D10P, K11H, E55A) (hereafter referred to as KPH and KPH-E55A, respectively) were prepared by chemical synthesis via ligation of 3 unprotected peptide segments, as described recently with minor modifications.16 One-pot chemical ligation was used to join the peptide fragments, which yielded the fully reduced 65-mer peptides. OD-KPH observed mass was 7219.2 ± 1.0 Da, calculated mass (average isotope composition) 7218.3 Da. OD1-KPH-E55A observed mass was 7160.1 ± 1.0 Da, calculated mass (average isotope composition) 7160.2 Da.

For in-vitro folding and disulfide bond formation, 10 mg (~1.4 μmol) of the purified and fully reduced peptides were dissolved in 6 M GdmHCl at a concentration of 4.0 mg/mL. Folding was initiated by rapid 1:20 dilution of the peptide solution with folding buffer (50 mM sodium phosphate, 8 mM reduced glutathione (GSH), 1mM oxidized glutathione (GSSG), pH 7.5). The reaction was left stirring at room temperature for 4 hours. The mixture was filtered and purified by HPLC on a Zorbax C3 column (10x250, 3 μm, 300 Å). OD1 KPH observed mass was 7210.5 ± 1.0 Da, calculated mass (average isotope composition) was 7210.3 Da. Yield 3.9 mg (0.6 μmol, 42%). OD1-KPH-E55A observed mass was 7153.2 ± 1.0 Da, calculated mass (average isotope composition) was 7152.2 Da. Yield: 3.2 mg (0.4 μmol, 29%).

**Electrophysiology**

CHO cells stably expressed with human Na$_{1.7}$ channels (Genionics AG, Zürich, Switzerland) were cultured as described previously and plated on round coverslips 1–4 d before experiments. Whole-cell Na$^+$ currents were recorded and filtered at 10 kHz using Axopatch 200B amplifier (Molecular Devices, Sunnyvale, CA) under voltage clamp conditions. Single-channel currents were filtered at 5kHz with a 24 dB/octave filter (Ithaco). The currents were digitized at 100 kHz using a Digidata 1400 interface and analyzed using pCLAMP 10.3 software package (Molecular Devices). Recording electrodes
in whole-cell experiments were pulled from TG150TF borosilicate glass capillaries (Harvard Apparatus) and had resistance of 1.2–1.6 MΩ when filled with a solution containing (mM): 130 CsF, 10 CsCl, 10 EGTA, 1 MgCl₂ and 10 HEPES-CsOH, pH 7.2. The extracellular solution contained (mM): 140 NaCl, 5 KCl, 1 MgCl₂, 2 CaCl₂, 10 glucose and 10 HEPES-NaOH, pH 7.3. Membrane potential was corrected for a calculated liquid junction potential of 6.9 mV. A series resistance of 2.7 ± 0.6 MΩ was compensated by 85–90%. Capacitive and leakage currents were digitally subtracted using a P/8 protocol with hyperpolarizing conditioning pulses. Single-channel recordings were performed in a outside-out configuration. Recording electrodes were pulled from aluminosilicate glass capillaries and had a resistance of 6–10 MΩ when filled with an intracellular solution.

Data analysis

The inactivation rate was estimated by the ratio of the current measured at 1.5 ms after the peak (I₁/₅) to the peak current value (Iₚ). Toxin concentration–response relationships were defined as a function of I₁/₅/Iₚ against concentration, and were fitted with a logistic equation: \( \frac{I_{1/5}}{I_p} = R_0 + \frac{R_1 - R_0}{1 + \left( \frac{C}{C_0} \right)^{n}} \), where C is toxin concentration; R₀, R₁ are asymptotic values for I₁/₅/Iₚ ratio. Conductance was calculated as: \( G = \frac{I_p}{V-V_r} \), where \( V_r \) is the reversal potential estimated by fitting current–voltage relationships with the equation:

\[
I = \frac{g(V-V_r)}{1 + \exp(-V/V_{1/2})},
\]

where \( V_{1/2} \) is the potential at which current peak reaches its half-maximum value. Voltage-dependence of activation and inactivation data were fitted by a Boltzmann function: \( f(V) = \frac{1}{1 + \exp(-V/V_{1/2})} \). Single-channel amplitude was estimated by fitting the amplitude histogram of elementary events with Gauss distribution. All non-linear regression analysis was performed using SigmaPlot 8.0 software. Results are shown as the mean ± standard error of the mean.

Kinetic modeling

To find the parameters of a kinetic model that give the best fit to whole-cell experimental current traces, we used a custom-developed application written in a MATLAB R2007b environment. In this approach, we resolved the optimization problem for the function

\[
\delta(K) = \sqrt{\sum_{i=1}^{n} (s_i(K) - y_i)^2},
\]

where K is the matrix of transition rates, \{s_i\} is a vector of n current samples derived from model equations and \{y_i\} is vector of sampled experimental whole-cell current. The problem was resolved by using a genetic algorithm minimizer with constraints – \( ga \), and/or gradient descent minimizer with constraints – \( fmincon \) (MATLAB). Differential equations associated with a kinetic model were numerically solved using \texttt{ode45} solver (MATLAB).

Disclosure of potential conflicts of interest

No potential conflicts of interest were disclosed.

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