

A Mutation in the Intracellular Loop III/IV of Mosquito Sodium Channel Synergizes the Effect of Mutations in Helix IIS6 on Pyrethroid Resistance[§]

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ABSTRACT

Activation and inactivation of voltage-gated sodium channels are critical for proper electrical signaling in excitable cells. Pyrethroid insecticides promote activation and inhibit inactivation of sodium channels, resulting in prolonged opening of sodium channels. They preferably bind to the open state of the sodium channel by interacting with two distinct receptor sites, pyrethroid receptor sites PyR1 and PyR2, formed by the interfaces of domains II/III and I/II, respectively. Specific mutations in PyR1 or PyR2 confer pyrethroid resistance in various arthropod pests and disease vectors. Recently, a unique mutation, N¹⁵⁷⁵Y, in the cytoplasmic loop linking domains III and IV (LIII/IV) was found to coexist with a PyR2 mutation, L¹⁰¹⁴F in IIS6, in pyrethroid-resistant populations of *Anopheles gambiae*. To ex-

amine the role of this mutation in pyrethroid resistance, N¹⁵⁷⁵Y alone or N¹⁵⁷⁵Y + L¹⁰¹⁴F were introduced into an *Aedes aegypti* sodium channel, AaNa_v1-1, and the mutants were functionally examined in *Xenopus* oocytes. N¹⁵⁷⁵Y did not alter AaNa_v1-1 sensitivity to pyrethroids. However, the N¹⁵⁷⁵Y + L¹⁰¹⁴F double mutant was more resistant to pyrethroids than the L¹⁰¹⁴F mutant channel. Further mutational analysis showed that N¹⁵⁷⁵Y could also synergize the effect of L¹⁰¹⁴S/W, but not L¹⁰¹⁴G or other pyrethroid-resistant mutations in IS6 or IIS6. Computer modeling predicts that N¹⁵⁷⁵Y allosterically alters PyR2 via a small shift of IIS6. Our findings provide the molecular basis for the coexistence of N¹⁵⁷⁵Y with L¹⁰¹⁴F in pyrethroid resistance, and suggest an allosteric interaction between IIS6 and LIII/IV in the sodium channel.

Introduction

Voltage-gated sodium channels are responsible for the rapidly rising phase of action potentials (Catterall, 2012). Because of their critical role in membrane excitability, sodium channels are the primary target site of a variety of naturally occurring and synthetic neurotoxins, including pyrethroid insecticides (Catterall et al., 2007). Pyrethroids promote activation and inhibit inactivation of sodium channels, resulting in prolonged opening of sodium channels (Vijverberg et al., 1982; Narahashi, 1996). Pyrethroid insecticides possess high insecticidal activities and low mammalian toxicity and represent one of the most powerful weapons in the global fight against malaria and other arthropod-borne human diseases. However, the efficacy of pyrethroids is undermined as a result of emerging pyrethroid resistance in arthropod pests and disease vectors. One major resistance mechanism is known as

knockdown resistance (kdr), which arises from mutations in the sodium channel (Soderlund, 2005; Rinkevich et al., 2013; Dong et al., 2014).

The pore-forming α -subunit of the sodium channel is composed by four homologous domains (I–IV), each having six transmembrane segments (S1–S6) connected by intracellular and extracellular loops. The S1–S4 segments in each domain serve as the voltage-sensing module, whereas the S5 and S6 segments and the loops connecting them function as the pore-forming module. In response to membrane depolarization, the S4 segments move outward, initiating conformational changes that lead to pore opening and subsequent inactivation of sodium channels. Short intracellular linkers connecting S4 and S5 segments of sodium channels, L45, transmit the movements of the voltage-sensing modules to the S6 segments during channel opening and closing. Fast inactivation is achieved by the movement of an inactivation gate formed mainly by the IFM motif in the short intracellular linker connecting domains III and IV, which physically occludes the open pore.

In recent years, using X-ray structures of a bacterial potassium channel KcsA (Doyle et al., 1998), the mammalian voltage-gated potassium channel K_v1.2 crystallized in the open state (Long et al., 2005) and a bacterial sodium channel,

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ABBREVIATIONS: AaNa_v, mosquito sodium channel; kdr, knockdown resistance; LIII/IV, cytoplasmic loop linking domains III and IV; PyR, pyrethroid receptor site.

Na_vAb, crystallized in the closed state (Payandeh et al., 2011) as templates, homology models of eukaryotic sodium channels have been developed to predict binding sites of sodium channel neurotoxins (Lipkind and Fozzard, 2005; O'Reilly et al., 2006; Tikhonov and Zhorov, 2007, 2012; Du et al., 2013). Mutational analyses coupled with computer modeling show that pyrethroids bind to two analogous receptor sites. Pyrethroid receptor site 1 (PyR1) is formed by residues from helices IIL45, IIS5, and IIS6 (O'Reilly et al., 2006), whereas pyrethroid receptor site 2 (PyR2) is formed by residues from helices IL45, IS5, IS6, and IIS6 (Du et al., 2013). Binding of pyrethroid molecules at the two sites is believed to effectively trap the sodium channel in the open state, resulting in the prolonged opening of sodium channels (Du et al., 2013).

The most frequent *kdr* mutation in arthropod pests and disease vectors is a leucine to phenylalanine (L¹⁰¹⁴F in the house fly sodium channel) in IIS6, which is also known as L²¹¹⁶F using the nomenclature that is universal for sodium channels and other P-loop ion channels (Zhorov and Tikhonov, 2004; Du et al., 2013) (Fig. 1). The L²¹¹⁶⁽¹⁰¹⁴⁾F mutation has been detected in the malaria vector *Anopheles* mosquito species in many regions around the world (Martinez-Torres et al., 1998; Enayati et al., 2003; Karunaratne et al., 2007). Recently, a new sodium channel mutation N¹⁵⁷⁵Y was reported in the malaria mosquito, *An. gambiae*, in Africa (Jones et al., 2012). The N¹⁵⁷⁵Y mutation is located in the intracellular cytoplasmic loop connecting domains III and IV (LIII/IV) (Fig. 1). Intriguingly, the N¹⁵⁷⁵Y mutation was only found in conjunction with L²¹¹⁶⁽¹⁰¹⁴⁾F: no mosquito individuals were detected harboring only the N¹⁵⁷⁵Y mutation (Jones et al., 2012). Interestingly, pyrethroid bioassays indicate that mosquitoes carrying the double mutations L²¹¹⁶⁽¹⁰¹⁴⁾F + N¹⁵⁷⁵Y are more resistant to permethrin than mosquitoes carrying only the L²¹¹⁶⁽¹⁰¹⁴⁾F mutation (Jones et al., 2012). However, whether the N¹⁵⁷⁵Y mutation confers pyrethroid resistance has not been functionally confirmed yet. In this study, we conducted site-directed mutagenesis, functional analysis in *Xenopus* oocytes, and computer modeling to investigate the role of N¹⁵⁷⁵Y in pyrethroid resistance.

Materials and Methods

Site-Directed Mutagenesis. Because sodium channels from *An. gambiae* have not been successfully expressed in the *Xenopus* oocyte expression system for functional characterization, we used a mosquito sodium channel (AaNa_v1-1), from *Aedes aegypti* to generate all mutants used in this study. The *kdr* mutations that are explored in this study are located in regions that are highly conserved between sodium

channels from *An. gambiae* and *Ae. aegypti* (Supplemental Fig. 1). Site-directed mutagenesis was performed by polymerase chain reaction using Pfu Turbo DNA polymerase (Stratagene, La Jolla, CA). All mutagenesis results were confirmed by DNA sequencing.

Expression of AaNa_v Sodium Channels in *Xenopus* Oocytes. The procedures for oocyte preparation and cRNA injection are identical to those described previously (Tan et al., 2002b). For robust expression of AaNa_v1-1 sodium channels, cRNAs were coinjected into oocytes with *Ae. aegypti* *TipE* cRNA (1:1 ratio), which enhances the expression of sodium channels in oocytes.

Electrophysiological Recording and Analysis. The voltage dependence of activation and inactivation was measured using the two-electrode voltage clamp technique. Methods for two-electrode recording and data analysis were identical to those described previously (Tan et al., 2002a).

The voltage dependence of sodium channel conductance (*G*) was calculated by measuring the peak current at test potentials ranging from -80 to +65 mV in 5 mV increments and divided by $(V - V_{rev})$, where *V* is the test potential and *V*_{rev} is the reversal potential for sodium ions. Peak conductance values were normalized to the maximal peak conductance (*G*_{max}) and fitted with a two-state Boltzmann equation of the form:

$$G/G_{max} = [1 + \exp(V - V_{1/2})/k]^{-1}$$

in which *V* is the potential of the voltage pulse, *V*_{1/2} is the voltage for half-maximal activation, and *k* is the slope factor.

The voltage dependence of sodium channel inactivation was determined by using 100 millisecond inactivating prepulses ranging from -120 to 10 mV in 5 mV increments from a holding potential of -120 mV, followed by test pulses to -10 mV for 20 milliseconds. The peak current amplitude during the test depolarization was normalized to the maximum current amplitude and plotted as a function of the prepulse potential. Data were fitted with a two-state Boltzmann equation of the form:

$$I/I_{max} = \{1 + [\exp(V - V_{1/2})/k]\}^{-1}$$

in which *I* is the peak sodium current, *I*_{max} is the maximal current evoked, *V* is the potential of the voltage prepulse, *V*_{1/2} is the half-maximal voltage for inactivation, and *k* is the slope factor. Sodium current decay was analyzed by fitting the decaying part of current traces with a single exponential function.

Measurement of Tail Currents Induced by Pyrethroids. The method for application of pyrethroids in the recording system was identical to that described previously (Tan et al., 2002a). The effects of pyrethroids were measured 10 minutes after their application. The pyrethroid-induced tail current was recorded during a 100 pulse train of 5 millisecond step depolarizations from -120 to 0 mV with 5 millisecond interpulse intervals (Vais et al., 2000). The percentage of channels modified by pyrethroids was calculated using the following equation (see Tatebayashi and Narahashi, 1994):

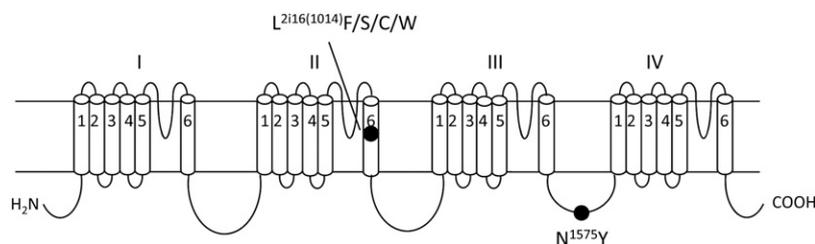


Fig. 1. The topology of the sodium channel protein indicating the position of L²¹¹⁶⁽¹⁰¹⁴⁾F/S/C/W and N¹⁵⁷⁵Y mutations. The sodium channel protein consists of four homologous domains (I–IV), each formed by six transmembrane segments (S1–S6) connected by intracellular and extracellular loops. Residue positions L¹⁰¹⁴F and N¹⁵⁷⁵Y correspond to the housefly sodium channel (GenBank accession numbers: AAB47604 and AAB47605). L²¹¹⁶F is labeled using the nomenclature that is universal for P-loop ion channels, in which a residue is labeled by the domain number (1–4), segment type (*k*, L45 linker; *i*, inner helix, i.e., S6; *o*, outer helix, i.e., S5), and the relative number of the residue in the segment.

$$M = \left\{ \frac{I_{\text{tail}} / (E_h - E_{\text{Na}})}{I_{\text{Na}} / (E_t - E_{\text{Na}})} \right\} \times 100$$

where I_{tail} is the maximal tail current amplitude, E_h is the potential to which the membrane is repolarized, E_{Na} is the reversal potential for sodium current determined from the current-voltage curve, I_{Na} is the amplitude of the peak current during depolarization before pyrethroid exposure, and E_t is the potential of step depolarization.

Molecular Modeling. We used the $K_v1.2$ -based model of the open AaNa_v1-1 channel with deltamethrin bound into PyR2 (Du et al., 2013) as a starting point to generate models of mutants with deltamethrin. Because pyrethroids preferably bind to sodium channels in the open state, the X-ray structure of the open potassium channel $K_v1.2$ (Long et al., 2005) was used as a template in this study and also in previous models of insect sodium channels with pyrethroids (O'Reilly et al., 2006, 2014; Du et al., 2013). The selectivity-filter region, which is substantially different between potassium and sodium channels, was not included in the models because it is far from the pyrethroid receptors (O'Reilly et al., 2006; Usherwood et al., 2007). Homology modeling and ligand docking were performed using the ZMM program (www.zmmsoft.com). Monte Carlo minimization protocol (Li and Scheraga, 1987) was used for energy optimization as described in Du et al. (2013) and Garden and Zhorov (2010). The complexes were visualized with the PyMol (Molecular Graphics System, Version 0.99rc6; Schrödinger, New York).

Chemicals. Deltamethrin was kindly provided by Bhupinder Khambay (Rothamsted Research, Harpenden, United Kingdom). Deltamethrin was dissolved in dimethylsulfoxide. The working concentration was prepared in ND96 recording solution immediately prior to experiments. The concentration of dimethylsulfoxide in the final solution was <0.5%, which had no effect on the function of sodium channels.

Statistical Analysis. Results are reported as mean \pm S.E.M. Statistical significance was determined by using one-way analysis of variance with Scheffe's post hoc analysis, and significant values were set at $P < 0.05$.

Results

$N^{1575}Y$ Does Not Alter the Gating Properties of AaNa_v1-1 Channels. The $N^{1575}Y$ mutation is located in LIII/IV, which is important for fast inactivation of sodium channels (Catterall, 2002; Goldin, 2003). To determine whether the $N^{1575}Y$ mutation alters inactivation kinetics, we introduced this mutation into AaNa_v1-1 and expressed both AaNa_v1-1 and mutant channels in *Xenopus* oocytes. As with AaNa_v1-1, the mutant channel produced sufficient sodium currents for functional analysis (Fig. 2). AaNa_v1-1 and $N^{1575}Y$ channels had similar rates of sodium current decay over the entire range of voltages examined, indicating that the mutation did not alter the kinetics of fast inactivation (Fig. 2). Furthermore, the mutation did not alter the voltage dependence of either activation or inactivation (Table 1).

Because the $N^{1575}Y$ mutation coexists with the $L^{216(1014)}F$ mutation in *An. gambiae* (Jones et al., 2012), the $N^{1575}Y$ change was also introduced into AaNa_v1-1 carrying the $L^{216(1014)}F$ mutation, which was generated in a previous study (Du et al., 2013), to create a double mutation construct $L^{216(1014)}F + N^{1575}Y$. Similarly, $L^{216(1014)}F$ and $L^{216(1014)}F + N^{1575}Y$ channels exhibited similar fast inactivation kinetics compared with AaNa_v1-1 (Fig. 2). The voltage dependences of activation and inactivation of both mutant channels were also similar to those of the AaNa_v1-1 channel (Table 1).

$N^{1575}Y$ Enhances $L^{216(1014)}F$ -Mediated Resistance to Pyrethroids, but Does Not Confer Pyrethroid Resistance Alone. To examine the effect of the $N^{1575}Y$ mutation on the

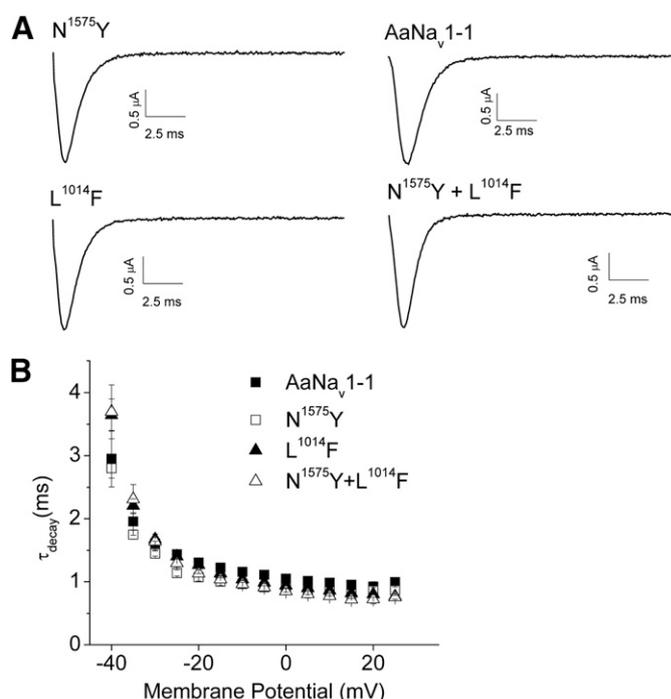


Fig. 2. $N^{1575}Y$ does not alter inactivation kinetics. (A) Representative current traces. The sodium currents were elicited by a 20-millisecond depolarization to 0 mV from the holding potential of -120 mV. (B) Voltage dependence of the sodium current decay. Time constant of sodium current decay was calculated by fitting the decaying part of current traces, elicited by a series of depolarizing voltages, with a single exponential function.

sensitivity of AaNa_v1-1 channels to pyrethroids, we compared the sensitivities of AaNa_v1-1, $N^{1575}Y$, $L^{216(1014)}F$, and $L^{216(1014)}F + N^{1575}Y$ channels to both a type I pyrethroid, permethrin, and a type II pyrethroid, deltamethrin. The percentage of sodium channel modification by pyrethroids was determined by measuring pyrethroid-induced tail currents upon repolarization in voltage-clamp experiments. The effect of permethrin on the $N^{1575}Y$ channel was similar to that on AaNa_v1-1 (Fig. 3, A and B). However, the permethrin-induced tail current was reduced by the $L^{216(1014)}F$ mutation and more drastically by $L^{216(1014)}F + N^{1575}Y$ double mutations (Fig. 3, B–D). Furthermore, analysis of the dose-response curves of modification of AaNa_v1-1 and the three mutant channels by permethrin and deltamethrin showed that the $L^{216(1014)}F$ mutant channel was about 8-fold more resistant to permethrin than the AaNa_v1-1 channel (Fig. 3E) and the $L^{216(1014)}F$ channel was 14-fold more resistant to deltamethrin than the AaNa_v1-1 channel (Fig. 3F). Remarkably, the $L^{216(1014)}F + N^{1575}Y$ channel was 80-fold more resistant to permethrin and 53-fold more resistant to deltamethrin than the wild-type channel. Therefore, the $N^{1575}Y$ mutation increased resistance to permethrin and deltamethrin by 9.8- and 3.4-fold, respectively, when combined with the $L^{216(1014)}F + N^{1575}Y$ mutation. However, the $N^{1575}Y$ mutation had no effect on channel sensitivity to either pyrethroid.

$N^{1575}Y$ Enhances Pyrethroid Resistance Caused by $L^{216(1014)}S/W$ Mutations. Our aforementioned results indicate that the $N^{1575}Y$ mutation imposes a synergistic effect on pyrethroid resistance caused by the $L^{216(1014)}F$ mutation. To see whether such synergism extends to pyrethroid resistance

TABLE 1

Voltage dependence of activation and fast inactivation of AaNa_v1-1 and its mutants.The voltage dependence of conductance and inactivation data were fitted with two-state Boltzmann equations, as described in the *Materials and Methods*, to determine $V_{1/2}$, the voltage for half-maximal conductance or inactivation and k , the slope for conductance or inactivation. Each value represents the mean \pm S.E.M.

	Activation		Fast Inactivation		
	$V_{1/2}$	k	$V_{1/2}$	k	n
	mV		mV		
AaNa _v 1-1	-33.6 \pm 1.1	5.7 \pm 0.3	-53.7 \pm 0.6	5.0 \pm 0.1	19
N ¹⁵⁷⁵ Y	-32.3 \pm 1.6	6.6 \pm 0.3	-56.5 \pm 0.5	5.0 \pm 0.1	20
L ²ⁱ¹⁶⁽¹⁰¹⁴⁾ F	-31.3 \pm 0.9	4.0 \pm 0.3	-49.6 \pm 0.4	4.5 \pm 0.1	10
L ²ⁱ¹⁶⁽¹⁰¹⁴⁾ F + N ¹⁵⁷⁵ Y	-28.6 \pm 1.1	5.6 \pm 0.2	-51.1 \pm 0.4	4.8 \pm 0.1	27
L ²ⁱ¹⁶⁽¹⁰¹⁴⁾ S	-36.3 \pm 1.1	4.3 \pm 0.3	-50.0 \pm 0.3	4.7 \pm 0.1	11
L ²ⁱ¹⁶⁽¹⁰¹⁴⁾ S + N ¹⁵⁷⁵ Y	-33.8 \pm 1.2	5.1 \pm 0.4	-51.5 \pm 0.7	4.9 \pm 0.1	10
L ²ⁱ¹⁶⁽¹⁰¹⁴⁾ W	-30.2 \pm 1.1	5.0 \pm 0.2	-50.3 \pm 0.5	4.8 \pm 0.1	6
L ²ⁱ¹⁶⁽¹⁰¹⁴⁾ W + N ¹⁵⁷⁵ Y	-29.8 \pm 1.2	5.2 \pm 0.2	-53.2 \pm 0.6	5.1 \pm 0.1	5
L ²ⁱ¹⁶⁽¹⁰¹⁴⁾ G	-31.5 \pm 1.0	4.9 \pm 0.3	-48.9 \pm 0.7	4.7 \pm 0.1	7
L ²ⁱ¹⁶⁽¹⁰¹⁴⁾ G + N ¹⁵⁷⁵ Y	-34.2 \pm 1.2	5.0 \pm 0.4	-52.3 \pm 0.6	4.6 \pm 0.1	9
L ^{2i16(L1014)} C	-37.5 \pm 1.1	6.2 \pm 0.3	-51.8 \pm 0.5	5.0 \pm 0.1	5
V ²ⁱ¹⁸⁽¹⁰¹⁶⁾ G	-33.0 \pm 1.0	5.3 \pm 0.3	-51.8 \pm 0.5	5.6 \pm 0.1	6
V ²ⁱ¹⁸⁽¹⁰¹⁶⁾ G + N ¹⁵⁷⁵ Y	-35.4 \pm 1.1	5.6 \pm 0.2	-52.1 \pm 0.7	5.3 \pm 0.1	7
L ¹ⁱ¹⁸ G	-30.1 \pm 1.0	4.8 \pm 0.5	-49.2 \pm 0.6	4.9 \pm 0.1	8
L ¹ⁱ¹⁸ G + N ¹⁵⁷⁵ Y	-32.5 \pm 1.2	4.9 \pm 0.4	-50.0 \pm 0.5	4.9 \pm 0.1	6
L ¹ⁱ¹⁸ F	-29.7 \pm 1.0	3.9 \pm 0.1	-42.9 \pm 0.7	5.0 \pm 0.1	8
L ¹ⁱ¹⁸ F + N ¹⁵⁷⁵ Y	-30.7 \pm 0.5	4.0 \pm 0.2	-44.2 \pm 0.6	5.2 \pm 0.1	7
L ¹ⁱ¹⁸ W	-29.0 \pm 1.1	4.2 \pm 0.2	-48.4 \pm 0.2	5.2 \pm 0.1	5
L ¹ⁱ¹⁸ W + N ¹⁵⁷⁵ Y	-28.2 \pm 0.6	4.8 \pm 0.2	-53.3 \pm 0.6	5.4 \pm 0.1	8
S ¹ⁱ²⁹ A	-33.2 \pm 1.0	5.2 \pm 0.3	-51.2 \pm 0.5	5.2 \pm 0.1	5
S ¹ⁱ²⁹ A + N ¹⁵⁷⁵ Y	-32.0 \pm 1.2	5.0 \pm 0.2	-50.2 \pm 0.6	5.1 \pm 0.1	6

caused by other *kdr* mutations identified in pyrethroid-resistant mosquitoes, we examined three additional *kdr* mutations, L²ⁱ¹⁶⁽¹⁰¹⁴⁾S/C/W, which have been reported in

various mosquito species (Ranson et al., 2000; Lüleyp et al., 2002; Stump et al., 2004; Kim et al., 2007; Kawada et al., 2009; Singh et al., 2010; Verhaeghen et al., 2010; Kasai et al., 2011;

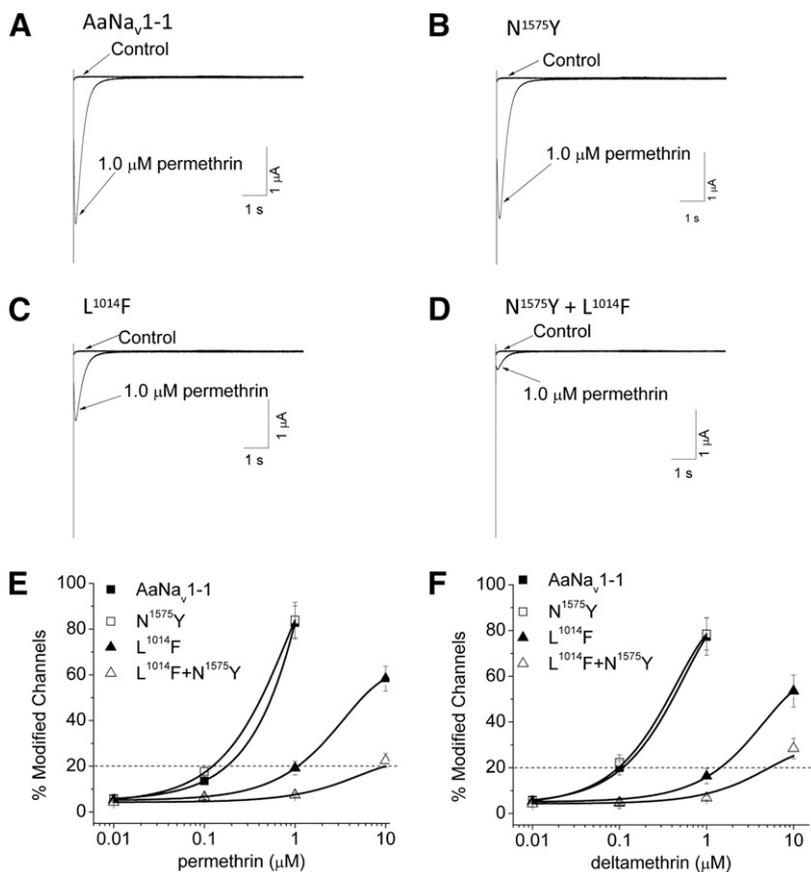


Fig. 3. The L¹⁰¹⁴F + N¹⁵⁷⁵Y double mutation reduced the sensitivity of AaNa_v1-1 channels to permethrin and deltamethrin. (A–D) Tail currents induced by permethrin (1 μ M) in AaNa_v1-1 (A), N¹⁵⁷⁵Y (B), L¹⁰¹⁴F (C), and L¹⁰¹⁴F + N¹⁵⁷⁵Y (D) channels. (E and F) Percentage of channel modification by permethrin (E) and deltamethrin (F) of wild-type and mutant channels. Percentage of channels modified by pyrethroids was calculated as described in the *Materials and Methods*. The values of EC₂₀ for permethrin were 0.12, 0.18, 1.0, and 9.8 μ M for AaNa_v1-1, N¹⁵⁷⁵Y, L¹⁰¹⁴F, and L¹⁰¹⁴F + N¹⁵⁷⁵Y channels, respectively. The values of EC₂₀ for deltamethrin were 0.1, 0.11, 1.4, and 5.3 μ M for AaNa_v1-1, N¹⁵⁷⁵Y, L¹⁰¹⁴F, and L¹⁰¹⁴F + N¹⁵⁷⁵Y channels, respectively. The number of oocytes for each mutant construct was >5. Each data point indicates mean \pm S.E.M. Asterisks indicate significant differences from the AaNa_v1-1 channel as determined by using one-way analysis of variance with Scheffe's post hoc analysis, and significant values were set at $P < 0.05$.

Tan et al., 2012; Wang et al., 2012). These mutations did not affect the voltage dependence of activation or inactivation of AaNa_v1-1 channels either alone or in conjunction with the N¹⁵⁷⁵Y mutation (Table 1). Consistent with results from a previous study (Du et al., 2013), the L²¹⁶⁽¹⁰¹⁴⁾S channel was 6.7- and 9-fold more resistant to permethrin and deltamethrin, respectively, than the AaNa_v1-1 channel (Fig. 4, A and B). The L²¹⁶⁽¹⁰¹⁴⁾S + N¹⁵⁷⁵Y double mutant channel was 58- and 32-fold more resistant to permethrin and deltamethrin, respectively, than the AaNa_v1-1 channel (Fig. 4, A and B). Similarly, the L²¹⁶⁽¹⁰¹⁴⁾W mutant channel was more resistant to permethrin and deltamethrin than the AaNa_v1-1 channel by about 9.7- and 7-fold, respectively (Fig. 4, C and D), and introduction of N¹⁵⁷⁵Y mutation into the L²¹⁶⁽¹⁰¹⁴⁾W channel caused additional 10.5- and 13-fold increase in permethrin and deltamethrin (101.8- and 91-fold resistance to permethrin and deltamethrin versus wild-type), respectively (Fig. 4, C and D). Another *kdr* mutation at the same position, L²¹⁶⁽¹⁰¹⁴⁾C, also caused 3.5- and 4.2-fold reduction in AaNa_v1-1 sensitivity to permethrin and deltamethrin, respectively (Supplemental Fig. 2). However, the L²¹⁶⁽¹⁰¹⁴⁾C + N¹⁵⁷⁵Y double mutant did not produce sufficient sodium currents in oocytes for further functional analysis.

N¹⁵⁷⁵Y Did Not Enhance Pyrethroid Resistance Caused by Another *kdr* Mutation in IIS6. Next, we examined another *kdr* mutation in IIS6, V²¹⁸⁽¹⁰¹⁶⁾G, which is detected in pyrethroid-resistant populations of *Ae. aegypti* (Saavedra-Rodriguez et al., 2007). Although all located in IIS6, L²¹⁶⁽¹⁰¹⁴⁾F/S/W and V²¹⁸⁽¹⁰¹⁶⁾G belong to PyR2 and PyR1, respectively (Du et al., 2013). We introduced the N¹⁵⁷⁵Y mutation into the V²¹⁸⁽¹⁰¹⁶⁾G channel construct, which was available from a previous study (Du et al., 2013). Consistent with the finding from the previous study, the V²¹⁸⁽¹⁰¹⁶⁾G mutation significantly reduced AaNa_v1-1 channel sensitivity to both permethrin and deltamethrin (Fig. 5, B and C), but the addition of the N¹⁵⁷⁵Y mutation did not further enhance the level of resistance to pyrethroids (Fig. 5, B and C).

N¹⁵⁷⁵Y Did Not Enhance Pyrethroid Resistance Caused by Other Mutations in PyR2. Many *kdr* mutations are located within the two pyrethroid-binding sites and/or are involved in regulating channel kinetics and voltage-dependent gating (Dong et al., 2014). N¹⁵⁷⁵Y is not located in either of the pyrethroid receptor sites and does not affect inactivation kinetics or channel gating (Fig. 2; Table 1). Therefore, we hypothesize that N¹⁵⁷⁵Y exerts its effect by allosterically altering one of the pyrethroid binding sites. The alteration per se must be small because N¹⁵⁷⁵Y alone does not affect the action of pyrethroids except in the presence of another resistance-associated mutation at L²¹⁶⁽¹⁰¹⁴⁾. Since the N¹⁵⁷⁵Y mutation enhances the effect of mutations at L²¹⁶⁽¹⁰¹⁴⁾, but not at V²¹⁸⁽¹⁰¹⁶⁾G, we then focused on residues in PyR2 to determine the extent of the N¹⁵⁷⁵Y-mediated synergism. In the PyR2 model (Du et al., 2013), the side chain of L²¹⁶⁽¹⁰¹⁴⁾ is directed toward the pyrethroid-sensing residue L¹¹⁸ in IS6 (Fig. 6A). Consistent with previous findings (Du et al., 2013), L¹¹⁸G reduced AaNa_v1-1 channel sensitivity to permethrin and deltamethrin. However, the double mutation L¹¹⁸G + N¹⁵⁷⁵Y was not more resistant to pyrethroids than the single mutant, L¹¹⁸G or S¹¹²⁹A (Fig. 5, B and C). Furthermore, we introduced two additional substitutions, L¹¹⁸F and L¹¹⁸W, into both the AaNav1-1 and N¹⁵⁷⁵Y channels. As with L¹¹⁸G, both substitutions reduced the action of pyrethroids (Fig. 5, B and C) and the N¹⁵⁷⁵Y mutation did not enhance pyrethroid resistance caused by either mutation (Fig. 5, B and C). In addition, S¹¹²⁹ at the C end of IS6 is predicted to approach the α -cyano group of deltamethrin and favorably contribute to ligand-channel interactions (Fig. 6, A and B). Here, we showed that S¹¹²⁹A decreased channel sensitivity to pyrethroids (Fig. 5, B and C), further supporting the PyR2 model. However, the S¹¹²⁹A + N¹⁵⁷⁵Y channel was not more resistant to both pyrethroids than the single S¹¹²⁹A mutant. Collectively, these results suggest that the modification of the receptor site by N¹⁵⁷⁵Y may be L²¹⁶⁽¹⁰¹⁴⁾ specific. It is possible that unfavorable interactions of the hydrophobic ligand with large aromatic substitutions L²¹⁶⁽¹⁰¹⁴⁾F/W or hydrophilic substitution L²¹⁶⁽¹⁰¹⁴⁾S

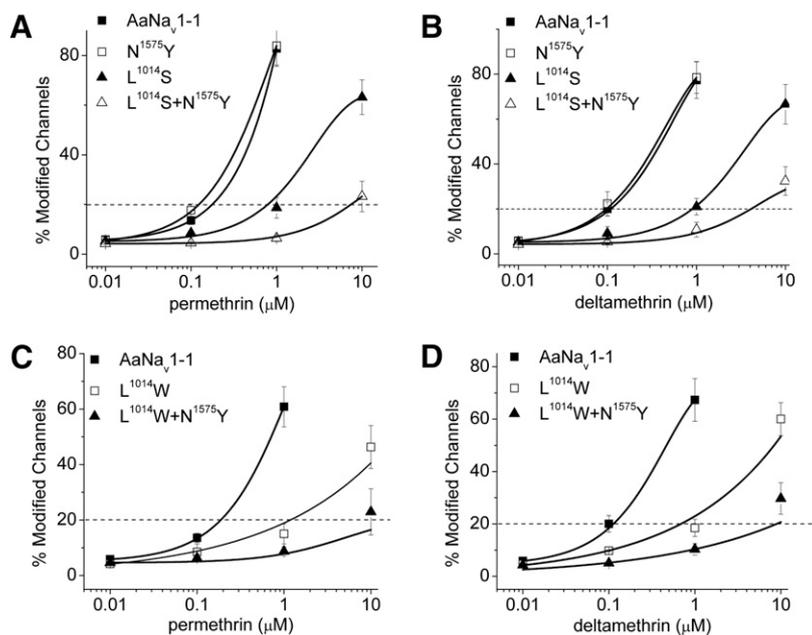


Fig. 4. N¹⁵⁷⁵Y enhanced resistance to pyrethroids caused by L1014S/W. Effects of L1014S/W (A, B and C, D, respectively) mutations on the sensitivity of AaNa_v1-1 channels to permethrin (A and C) or deltamethrin (B and D). Percentages of channel modification by permethrin or deltamethrin were determined using the method described in the *Materials and Methods*. The number of oocytes for each mutant construct was >5. Each data point indicates mean \pm S.E.M. Asterisks indicate significant differences from the AaNa_v1-1 channel as determined by using one-way analysis of variance with Scheffe's post hoc analysis, and significant values were set at $P < 0.05$.

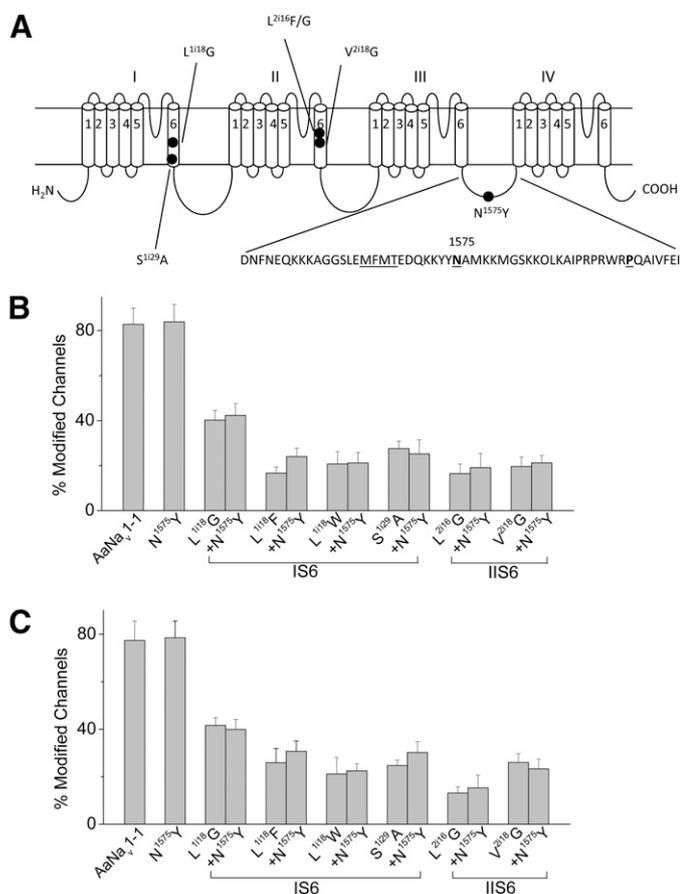


Fig. 5. $N^{1575}Y$ did not enhance pyrethroid resistance caused by other mutations examined in this study. (A) The positions of the mutations in the sodium channel protein. The amino acid sequence of the linker connecting domains III and IV is shown below the topology. The MFMT motif, which is critical for fast inactivation, is underlined. The $N^{1575}Y$ mutation is marked in bold. Residue P^{1596} in this linker, whose leucine substitution was previously confirmed to increase pyrethroid potency, is also shown in bold. (B and C) Effects of $N^{1575}Y$ on the sensitivity of $AaNa_v1-1$ wild-type and mutant channels to 1 μ M permethrin (B) and 1 μ M deltamethrin (C). The number of oocytes for each mutant construct was >5 . Each data point indicates mean \pm S.E.M. The percentage of channel modification for all mutant channels except $N^{1575}Y$ was significantly different from that of the $AaNa_v1-1$ channel, but with no significant difference between the double mutants and their respective singles, as determined by using one-way analysis of variance with Scheffé's post hoc analysis, and significant values were set at $P < 0.05$.

were enhanced by $N^{1575}Y$. Thus, we examined a glycine substitution of $L^{216(1014)}$, i.e., $L^{216(1014)}G$, for pyrethroid sensitivity. As with $L^{216(1014)}F/S/W$, the $L^{216(1014)}G$ substitution reduced the $AaNa_v1-1$ channel sensitivity to permethrin and deltamethrin. However, the $L^{216(1014)}G + N^{1575}Y$ double mutant channel was as sensitive as the $L^{216(1014)}G$ channel to both permethrin and deltamethrin (Fig. 5, B and C). These results suggest that the effect of $N^{1575}Y$ on pyrethroid binding, potentially causing a small conformational shift in IIS6, is sensitive to the nature of the residue at position 2i16 (1014).

Possible Mechanism of Synergy between $N^{1575}Y$ and $L^{216(1014)}F/S/W$ Mutations. To explore how $N^{1575}Y$ enhances pyrethroid resistance of $AaNa_v1-1$ channels when combined with the $L^{216(1014)}F/S/W$ mutations, but not in wild-type or with the $L^{216}G$ mutation, we compared the model of deltamethrin binding in PyR2 of the wild-type open $AaNa_v1-1$ channel and the mutants (Fig. 6). PDB format files for the $AaNa_v1-1$ channel

models are provided in the Supplemental Material. In the wild-type channel, deltamethrin favorably interacts with the long flexible side chain of $L^{216(1014)}$ (Fig. 6A). Comparison of deltamethrin binding models in the wild-type channel and mutants $L^{118}G$ (Fig. 6C) and $L^{216}G$ (Fig. 6D) suggests two consequences of the glycine substitutions. First, the favorable interactions of the hydrophobic ligand with the large hydrophobic pyrethroid-sensing residues are lost. Second, there is significantly more space in the pyrethroid binding site following mutation of L^{216} to G. Models of deltamethrin binding in the $L^{216}S$ and $L^{216}F$ mutants show unfavorable interactions between the hydrophilic S^{216} or the large and inflexible F^{216} with the hydrophobic ligand (Fig. 6, E and F). Therefore, we suggest that enhancement of pyrethroid resistance in the presence of the $L^{216(1014)}F/S/W$ mutations occurs as a result of a slight shift of IIS6. The shift does not affect either the favorable interaction of pyrethroids with L^{216} or the less sterically constrained interactions of deltamethrin with the $L^{216}G$ mutation. In contrast, the slight shift of IIS6 likely further deteriorates the energetically unfavorable interactions of deltamethrin with $L^{216(1014)}F/S/W$, enhancing pyrethroid resistance in the double mutant.

Discussion

Identification of naturally occurring mutations in the sodium channel that confer kdr to pyrethroids has greatly advanced our understanding of the molecular mechanisms of kdr and the molecular details of pyrethroid receptor sites (Dong et al., 2014). Emerging evidence suggests that binding of pyrethroids to two distinct pyrethroid receptor sites, PyR1 and PyR2, at two analogous domain interfaces is necessary to trap sodium channels in the open state, which leads to prolonged opening of sodium channels and the toxic effects of pyrethroids in vivo. Importantly, many, but not all, kdr mutations that cause pyrethroid resistance are located within the two pyrethroid receptor sites (Dong et al., 2014), and it is not clear how kdr mutations beyond the receptor sites affect pyrethroid sensitivity of sodium channels. In this study, we investigated the role of a pyrethroid resistance-associated mutation, $N^{1575}Y$, which is located in the intracellular loop connecting domains III and IV (outside of PyR1 and PyR2) in pyrethroid resistance. We found that this mutation alone has no effect on the action of pyrethroids on sodium channels, but it enhances pyrethroid resistance caused by the $L^{216(1014)}F/S/W$ mutations. Based on further mutational analysis and computer modeling, we hypothesize that $N^{1575}Y$ induces a small shift of the transmembrane helix IIS6, resulting in slight deformation of PyR2, which enhances the energetically unfavorable interactions between deltamethrin and the $L^{216(1014)}F/S/W$ mutations, but does not affect the energetically favorable interactions between pyrethroids and $L^{216(1014)}$. These results provide a satisfying explanation for the concurrent existence of $N^{1575}Y$ and $L^{216(1014)}F$ in pyrethroid-resistant mosquito populations.

BLAST searches show that the asparagine residue (equivalent to N^{1575} in *An. gambiae*) in the LIII/IV loop sequence is highly conserved among voltage-gated sodium channels. However, $N^{1575}Y$ has no effect on the kinetics of fast inactivation, implying that this mutation does not interfere with the docking of the inactivation particle (the motif MFMT in LIII/IV) to its receptor site, which is composed of residues in the linkers connecting S4 and S5 of domains III and IV (Goldin, 2003). Our

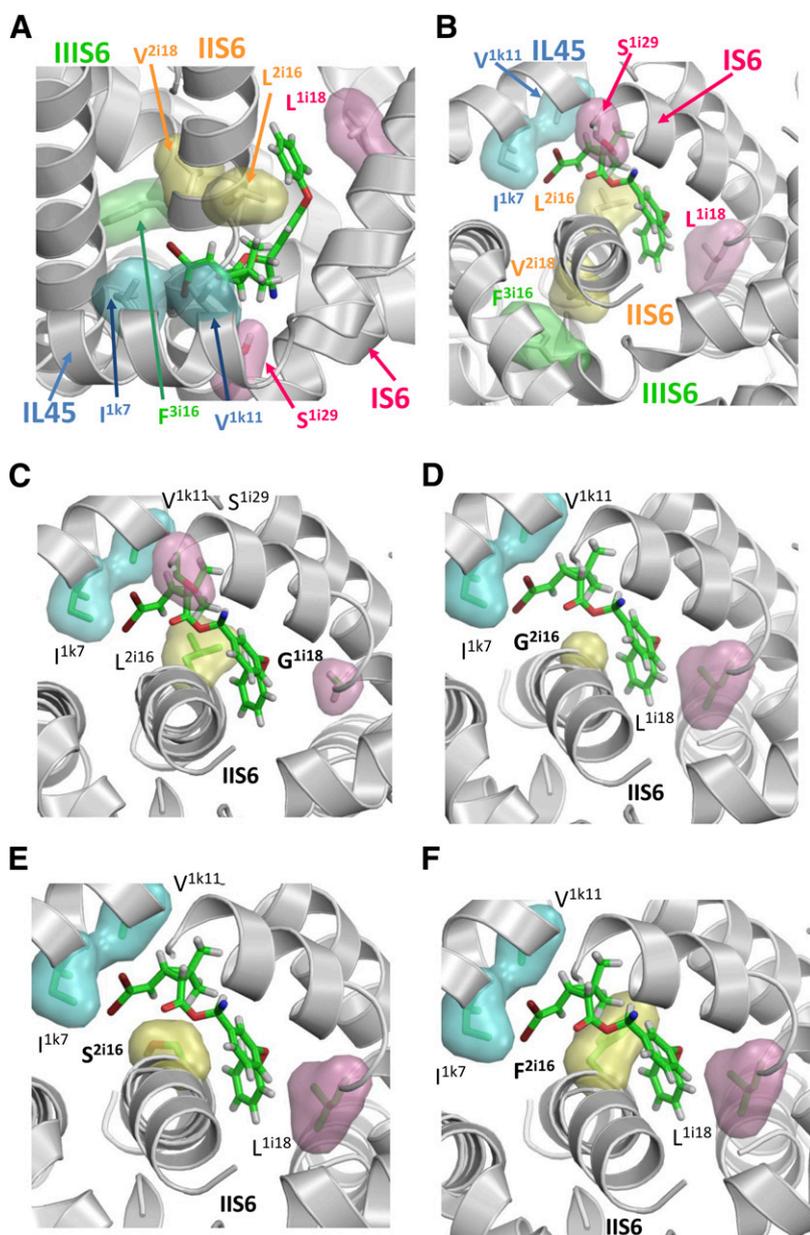


Fig. 6. K_v1.2-based model of the open AaNa_v1-1 channel with deltamethrin bound to PyR2. Side (A) and cytoplasmic (B) views. Deltamethrin is shown by sticks with green carbons, red oxygens, gray hydrogens, and brown bromine atoms. Semitransparent cyan, pink, yellow, and green surfaces show pyrethroid-sensing residues in helices IL45, IS6, IIS6, and IIS6, respectively. Note that helix IIS6 contributes residue L²¹⁶ to PyR2 and may contribute residue V²¹⁸ to PyR1, which contains F³¹⁶ in helix IIS6. (C–F), Cytoplasmic view along helix IIS6 of models of AaNa_v1-1 mutants L¹¹⁸G (C), L²¹⁶G (D), L²¹⁶S (E), and L²¹⁶F (F) of the open AaNa_v1-1 channel with deltamethrin bound in the pyrethroid receptor PyR2. The long flexible side chain of L²¹⁶ in the L¹¹⁸G mutant (C) and the wild-type channel (B) favorably interact with the ligand and a small shift of helix IIS6 upon mutation N¹⁵⁷⁵Y would have little effect on this interaction. The ligand potency in the L²¹⁶G mutant (D) would also have low sensitivity to the IIS6 shift because the tiny side chain of G²¹⁶ is located rather far from the ligand. However, the IIS6 shift would deteriorate unfavorable interactions of the hydrophilic S²¹⁶ with the hydrophobic ligand (E) or would cause an unfavorable clash of deltamethrin with the big and inflexible F²¹⁶ (D).

molecular modeling (Fig. 6) predicts that a slight shift of helix IIS6 in the N¹⁵⁷⁵Y channel could explain the synergism on pyrethroid resistance between N¹⁵⁷⁵Y and specific mutations at L²¹⁶⁽¹⁰¹⁴⁾F in IIS6, which is also supported by our mutational analysis (Figs. 3–5). However, we cannot completely rule out the possibility of the N¹⁵⁷⁵Y/L²¹⁶⁽¹⁰¹⁴⁾F double mutation causing an allosteric effect on the action of pyrethroids without directly involving IIS6.

How a mutation in the intracellular loop between domains III and IV (LIII/IV) can shift helix IIS6 carrying the L²¹⁶⁽¹⁰¹⁴⁾F mutation remains speculative. In X-ray structures of both open and closed ion channels, the cytoplasmic parts of the S6 helices approach each other to form the activation gate, and cytoplasmic linkers may also approach each other. We speculate that the asparagine side chain in LIII/IV is involved in specific contacts (likely, an H-bond) with LII/III. Replacement of a small asparagine with a much bigger tyrosine in N¹⁵⁷⁵Y could cause a change in the mutual disposition of the two linkers, which, in

turn, would shift helix IIS6. This predicted small shift in IIS6 by N¹⁵⁷⁵Y apparently only alters the action of pyrethroids on L²¹⁶⁽¹⁰¹⁴⁾F/S/C/W channels, but not on L²¹⁶G and V²¹⁸⁽¹⁰¹⁶⁾G channels. This may be because a small distortion of PyR2 upon the N¹⁵⁷⁵Y mutation does not affect a weak pyrethroid interaction with a glycine residue, which has a single hydrogen atom in the side chain.

Besides N¹⁵⁷⁵Y, there are several other sodium channel mutations within LIII/IV that have been reported to be associated with pyrethroid resistance (Rinkevich et al., 2013; Dong et al., 2014). The frequent occurrence of mutations associated with pyrethroid resistance in this linker supports the allosteric interactions between IIS6 and LIII/IV in the sodium channel. For example, L¹⁵⁹⁶P was found to be associated with pyrethroid resistance in *Varroa* mites (Fig. 5) (Dong et al., 2014). Although this mutation has not been functionally examined using *Varroa* mite sodium channels expressed in *Xenopus* oocytes, insect sodium channels possess

a proline at the corresponding position and the leucine substitution of proline in the cockroach sodium channel renders the cockroach sodium channel more sensitive to pyrethroids (Liu et al., 2006). Therefore, the L¹⁵⁹⁶P mutation is predicted to make the *Varroa* mite sodium channel more resistant to pyrethroids. Since L¹⁵⁹⁶P alone could confer pyrethroid resistance, P1596, located at the C-terminus of LIII/IV, likely has a more drastic allosteric effect on pyrethroid binding (to PyR1 and/or PyR2) than N¹⁵⁷⁵Y.

The involvement of other intracellular linkers in pyrethroid resistance has also been reported. E⁴³⁵K and C⁷⁸⁵R, in the linker connecting domains I and II (LI/II), were identified in pyrethroid resistant German cockroach populations (Liu et al., 2000; Dong et al., 2014). As with N¹⁵⁷⁵Y, E⁴³⁵K or C⁷⁸⁵R alone did not reduce sodium channel sensitivity to pyrethroids. However, concurrence of either E⁴³⁵K or C⁷⁸⁵R mutation with the *kdr* mutations, V¹¹¹⁹⁽⁴¹⁰⁾M in IS6 or L²¹⁶⁽¹⁰¹⁴⁾F in IIS6, significantly increases pyrethroid resistance (Liu et al., 2002; Tan et al., 2002b). A mechanism similar to that for N¹⁵⁷⁵Y could explain the role of E⁴³⁵K or C⁷⁸⁵R in pyrethroid resistance. Another example is G¹¹¹¹ (in the cockroach sodium channel) in the second intracellular linker connecting domains II and III (LII/III), which is selectively involved in the response of sodium channels to type II pyrethroids, such as deltamethrin (Du et al., 2009). Deletion of G¹¹¹¹ (due to alternative splicing of cockroach sodium channel transcripts) makes cockroach sodium channels more resistant to type II pyrethroids. Interestingly, although the overall sequence of the intracellular linker is quite variable, the amino acid sequence around G¹¹¹¹ is highly conserved among insect sodium channels. Two conserved lysine residues K¹¹¹⁸ and K¹¹¹⁹ downstream from G¹¹¹¹ are also critical for the action of type II pyrethroids (Du et al., 2009). Neutralization of K¹¹¹⁸ and K¹¹¹⁹ confers resistance to type II pyrethroids. The precise mechanism through which these mutations selectively alter the interaction of sodium channels with pyrethroids remains unclear. It is possible that they alter the binding site for type II pyrethroids by an allosteric mechanism.

In conclusion, our study sheds light on the mechanism by which the N¹⁵⁷⁵Y mutation enhances pyrethroid resistance and explains the molecular basis of the concurrence of N¹⁵⁷⁵Y and L²¹⁶⁽¹⁰¹⁴⁾F in pyrethroid-resistant mosquito populations. Furthermore, our findings provide evidence for possible allosteric, cross-domain interactions between transmembrane segments and intracellular loops of the sodium channel in mediating the action of pyrethroids on sodium channels.

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Authorship Contributions

Participated in research design: Wang, Zhorov, Dong.
Conducted experiments: Wang, Du, Nomura.
Performed data analysis: Wang, Du, Nomura.
Wrote or contributed to the writing of the manuscript: Wang, Du, Liu, Zhorov, Dong.

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