Localization and Molecular Determinants of the Hanatoxin Receptors on the Voltage-sensing Domains of a K⁺ Channel

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Abstract Hanatoxin inhibits voltage-gated K⁺ channels by modifying the energetics of activation. We studied the molecular determinants and physical location of the Hanatoxin receptors on the drk1 voltage-gated K⁺ channel. First, we made multiple substitutions at three previously identified positions in the COOH terminus of S3 to examine whether these residues interact intimately with the toxin. We also examined a region encompassing S1–S3 using alanine-scanning mutagenesis to identify additional determinants of the toxin receptors. Finally, guided by the structure of the KcsA K⁺ channel, we explored whether the toxin interacts with the peripheral extracellular surface of the pore domain in the drk1 K⁺ channel. Our results argue for an intimate interaction between the toxin and the COOH terminus of S3 and suggest that the Hanatoxin receptors are confined within the voltage-sensing domains of the channel, at least 20–25 Å away from the central pore axis.

Key words: gating modifier toxin • scanning mutagenesis • voltage-dependent gating • protein–protein interaction

Introduction

Voltage-gated K⁺ channels are tetramers with each subunit containing six transmembrane segments, termed S1–S6. As illustrated in Fig. 1, we can think of these channels as constructed from two types of domains: a central pore domain formed by S5–S6 and four surrounding voltage-sensing domains, each comprised of S1–S4. The structure of the pore domain is likely to be similar to the crystal structure of the KcsA K⁺ channel, a bacterial channel that is homologous to S5–S6 in voltage-gated K⁺ channels (Schrempf et al., 1995; Doyle et al., 1998; MacKinnon et al., 1998; Heginbotham et al., 1999). Thus, both S5 and S6 in voltage-gated K⁺ channels are membrane-spanning α helices with the linker between them forming a short pore helix and the selectivity filter. A growing body of evidence suggests that the voltage-sensing domains are constructed from S1 through S4 (Papazian et al., 1991, 1995; Liman et al., 1991; Perozo et al., 1994; Planells-Cases et al., 1995; Aggarwal and MacKinnon, 1996; Larsson et al., 1996; Mannuzzu et al., 1996; Seoh et al., 1996; Yang et al., 1996; Yusaf et al., 1996; Tiwari-Woodruff et al., 1997; Cha and Bezanilla, 1997; Smith-Maxwell et al., 1998a,b; Li-Smerin and Swartz, 1998; Ledwell and Aldrich, 1999; Monks et al., 1999; Hong and Miller, 2000; Li-Smerin et al., 2000a). Compared with the pore domain, much less is known about the structure of this important region of voltage-gated K⁺ channels. As illustrated in Fig. 1 A, the first four transmembrane segments (S1, S2, S3, and S4) are thought to be membrane-spanning α helices (Monks et al., 1999; Hong and Miller, 2000; Li-Smerin et al., 2000a). There is also evidence for additional helices in the extracellular linkers between S1 and S2 and between S3 and S4 (Li-Smerin et al., 2000a).

A large number of protein toxins isolated from venomous animals interact with voltage-gated ion channels and either physically block ion conduction or modify voltage-dependent gating. The pore-blocking toxins interact with the outer vestibule of the channel, as illustrated by the receptor for Agitoxin, in Fig. 1 B (MacKinnon and Miller, 1988, 1989; Miller, 1988, 1995; Hurst et al., 1991; Stocker et al., 1991; Park and Miller, 1992a,b; Goldstein and Miller, 1993; Garcia et al., 1994; Goldstein et al., 1994; Lu and MacKinnon, 1997; Imredy et al., 1998; Jin and Lu, 1998). In fact, these toxins were initially used to identify the pore-forming region of the channel and subsequently to probe the three-dimensional structure of this region (MacKinnon and Miller, 1988; Hurst et al., 1991; Stampe et al., 1994; Stocker and Miller, 1994; Goldstein et al., 1994; Hidalgo and MacKinnon, 1995; Ranganathan et al., 1996; Naranjo and Miller, 1996; MacKinnon et al., 1998). Importantly, many of the structural constraints obtained using pore-blocking toxins have been born out by the crystal structure of the KcsA K⁺ channel, recently solved by x-ray diffraction (Doyle et al., 1998; MacKinnon et al., 1998). In contrast to the pore-blocking toxins, the location and structural determinants of the receptors for gating modifier toxins are poorly un-
Hanatoxin is a protein toxin from spider venom that inhibits the drk1 voltage-gated K\(^+\) channel by binding to receptors on the extracellular face of the channel and modifying the energetics of gating (Swartz and MacKinnon, 1995, 1997a). This mechanism was suggested by the observation that toxin-bound channels can open, but require large depolarizing voltages to do so. Unlike the pore-blocking toxins that bind with 1:1 stoichiometry, four Hanatoxin molecules can bind to the external surface of a single K\(^+\) channel (Swartz and MacKinnon, 1997a). An initial search for residues forming the Hanatoxin receptors identified three positions (I273, F274, and E277), where point mutations have been found to alter the binding affinity of Hanatoxin (Swartz and MacKinnon, 1997b). The mutative demarcations of the S3 and S4 transmembrane segments, indicated by underlining, are based on Kyte-Doolittle hydrophobicity analysis. The cylinders above the sequence indicated the approximate positions of three \(\alpha\) helices. The solid cylinders (for S3 and S4) are based on \(\alpha\)-helical periodicity detected using alanine- and tryptophan-scanning mutagenesis (Hong and Miller, 2000; Li-Smerin et al., 2000a). The dotted cylinder positioned near the COOH terminus of S3 and the S3–S4 linker was proposed based on helical periodicity detected with alanine-scanning mutagenesis (Li-Smerin et al., 2000a). The topology of a single K\(^+\) channel subunit shown (bottom) illustrates the first four transmembrane segments (S1–S4, in black) comprising the voltage-sensing domain and the last two transmembrane segments (S5 to S6, in blue) the pore domain.

The dotted cylinder in the S1–S2 linker is based on helical periodicity detected with alanine-scanning mutagenesis (Li-Smerin et al., 2000a). The top is extracellular and the bottom is intracellular. (B) Architecture of voltage-gated K\(^+\) channels containing a central pore domain and four surrounding voltage-sensing domains drawn according to Li-Smerin et al. (2000b). The pore domain is represented by the crystal structure of KcsA (Doyle et al., 1998), a simple K\(^+\) channel homologous to the S5–S6 region of voltage-gated K\(^+\) channels. Adjacent pore domain subunits in the tetramer are illustrated using different shades of blue. The red ring over the ion conduction pore indicates the footprint of Agitoxin\(_2\) (\(\sim\)20 \(\times\) 30 Å), a pore-blocking toxin (MacKinnon et al., 1998). Each surrounding voltage-sensing domain is formed by S1–S4 transmembrane helices. The four areas surrounded by black dotted lines are regions that potentially interact with Hanatoxin, based on cohabitation of Hanatoxin and Agitoxin\(_2\). The overall dimensions of the channel are drawn using estimates of 80 \(\times\) 80 Å from electron microscopy (Li et al., 1994).
with Hanatoxin (Fig. 1) (Swartz and MacKinnon, 1997b). The cohabitation of pore-blocking and gating-modifier toxins argues for peripheral locations of the Hanatoxin receptors, probably >15 Å from the central pore axis. The peripheral location of the Hanatoxin receptors fits nicely with the 4:1 stoichiometry between toxin and channel since these channels are homotetramers. However, the actual position of the Hanatoxin receptors remains poorly defined because the pore-blocking toxin receptor constitutes a relatively small fraction of the large (~80 × 80 Å) extracellular surface of the channel (Li et al., 1994). Thus, Hanatoxin might interact with the channel anywhere within the black dotted lines in Fig. 1 B.

In this paper, we studied the structural components and location of the Hanatoxin receptors on the drk1 K+ channel. First, we examined whether the three previously identified residues in the COOH-terminal end of S3 interact with the toxin by looking for patterns arising from multiple substitutions at each position. Second, we examined the possible contribution of additional residues in forming the Hanatoxin receptors by Ala-scanning a previously unexplored region between the NH2-terminal side of S1 and the COOH-terminal end of S3. Finally, we explored the possibility that Hanatoxin interacts with residues on the peripheral surface of the pore domain. Our results are consistent with an intimate interaction between Hanatoxin and channel residues in the COOH-terminal part of S3. In addition, our results suggest that the toxin does not interact with the extracellular surface of the pore domain, but rather exclusively interacts with the voltage-sensing domains of the channel.

**M A T E R I A L S A N D M E T H O D S**

**Mutagenesis and Channel Expression**

The cDNA encoding the wild-type drk1 K+ channel (Frech et al., 1989) used in this study contained several unique restriction sites that were previously introduced using PCR (Swartz and MacKinnon, 1997b). To construct point mutations of the channel, mutagenic residues in the COOH-terminal end of S3 interact with the toxin by looking for patterns arising from multiple substitutions at each position. Mutations were confirmed by dideoxy (Sanger et al., 1977) or automated DNA sequencing. cDNAs encoding wild-type and mutant channels were linearized with NotI and transcribed with T7 RNA polymerase.

Oocytes from *Xenopus laevis* frogs were removed surgically and incubated with agitation for 1-1.5 h in a solution containing (mM): 82.5 NaCl, 2.5 KCl, 1 MgCl2, 5 HEPES, and 2 mg/ml collagenase (Worthington Biochemical Corp.), pH 7.6 with NaOH. Defolliculated oocytes were injected with cRNA and incubated at 17°C in a solution containing (mM): 50 NaCl, 2 KCl, 1 MgCl2, 1.8 CaCl2, 5 HEPES, 50 μg/ml gentamicin (GIBCO BRL), pH 7.6 with NaOH for 1-5 d before recording.

**Examination of the Toxin-Channel Interaction**

Two-electrode voltage-clamp recording techniques were employed to study the toxin-channel interaction using an OC-725C oocyte clamp (Warner Instruments). Oocytes were studied in a 160-μl recording chamber that was perfused with a solution containing (mM): 50 RbCl, 50 NaCl, 1 MgCl2, 0.3 CaCl2, and 5 HEPES, pH 7.6 with NaOH. Data were filtered at 2 kHz (eight-pole Bessel) and digitized at 10 kHz. Microelectrode resistances were between 0.2 and 1.2 MΩ when filled with 3 M KCl. All experiments were performed at room temperature (~22°C).

The equilibrium dissociation constant for Hanatoxin binding to closed or resting channels was determined using very negative holding voltages (~120 to ~80 mV) where no steady state inactivation could be detected. The fraction of unbound channels was estimated using depolarizations that were too weak to open toxin bound channels and too short to perturb the equilibrium for toxin binding to resting channels, as previously described (Swartz and MacKinnon, 1997a,b). After toxin-bound channels are opened by strong depolarization, they close more rapidly than unbound channels after repolarization (Swartz and MacKinnon, 1997a). We therefore examined the kinetics of deactivation using tail currents carried by 50 mM Rb+ (to slow deactivation) to confirm that toxin-bound channels did not contribute to the currents measured with weak depolarizations. For all channels, we recorded tail current voltage-activation relations in the absence and presence of different concentrations of toxin. After addition of the toxin to the recording chamber, the equilibrium between toxin and channel was monitored using weak depolarizations elicited at 5-204 intervals. Recovery from toxin inhibition after removing the toxin from the recording chamber was routinely monitored but not examined quantitatively because of complexities arising from the 4:1 stoichiometry between toxin and channel (Swartz and MacKinnon, 1997a,b). The ratio of tail current (I/I0) recorded in the presence (I) and absence (I0) of toxin was calculated for various strength depolarizations, typically ~50 to ~60 mV. The value of I/I0 measured in the plateau phase at negative voltages (I/I0), where toxin-bound channels do not open, was taken as equal to the fraction of unbound channels (see Fig. 2). The equilibrium dissociation constant (Kd) for toxin binding was calculated according to:

\[
K_d = \frac{1}{1 - \left( \frac{I}{I_0} \right)}
\]

This equation assumes four independent and equivalent binding sites on the drk1 K+ channel for Hanatoxin. For mutant channels with altered gating properties, all voltage protocols were adjusted appropriately so that the plateau phase in the I/I0-voltage relation was well defined. The fraction of uninhibited current (I/I0) was initially measured for all mutant channels using between 100 and 250 nM Hanatoxin. The concentration dependence for toxin inhibition was further examined using two or three toxin concentrations for all mutants displaying >10-fold changes in toxin Kd in the initial screen.

Hanatoxin was purified from tarantula venom as previously described (Swartz and MacKinnon, 1995). The native toxin preparation is an approximately equal mixture of two isoforms that differ only at position 13, where Hanatoxin1 contains serine and Hanatoxin2 contains alanine. The two isoforms most likely have similar binding affinities because the concentration dependence for inhibition of the channel by the native mixture and synthetic Hanatoxin1 are similar (Takahashi et al., 2000).

**R E S U L T S**

Hanatoxin inhibits the drk1 voltage-gated K+ channel by shifting activation to more depolarized voltages (Swartz and MacKinnon, 1997a). We set out to localize the Hanatoxin receptors on the drk1 voltage-gated K+ channel.
channel by examining the effects of point mutations on toxin binding affinity. Wild-type and mutant channels were expressed in *Xenopus* oocytes and the interaction between toxin and channel were examined electrophysiologically. We determined the equilibrium dissociation constant for toxin binding to mutant and wild-type channels by studying the concentration dependence for toxin occupancy of the channel as previously described (Swartz and MacKinnon, 1997a) (see materials and methods). In this approach, the fraction of unbound channels is estimated by measuring the fraction of uninhibited current at negative voltages where toxin bound channels do not open.

Effects of Multiple Substitutions in the COOH-terminal Part of S3

Mutations to Ala at three positions in the COOH-terminal part of S3 in the drk1 K⁺ channel were previously reported to decrease Hanatoxin binding affinity by ~10-25-fold (Swartz and MacKinnon, 1997b). In the wild-type channel, two of these residues are hydrophobic (I273 and F274) and one is acidic (E277). To explore possible interactions of these three channel residues with the toxin, we made multiple substitutions (one at a time) at each of these positions and determined the equilibrium dissociation constant for Hanatoxin binding to each mutant channel. Our approach was to look for patterns between changes in the toxin Kₐ and the chemical nature of the substituted amino acids.

The measurement of toxin binding affinity is illustrated in Fig. 2 for the wild-type and two exemplary mutant channels (F274R and E277K). Voltage-activation relations were obtained in the absence and presence of various toxin concentrations using tail-current protocols (Fig. 2A and B). The fraction of uninhibited tail current was then measured for different strength depolarizations, as shown in Fig. 2C. This fraction of uninhibited current in the plateau phase at negative voltages is plotted as a function of test voltage for wild-type, F274R, and E277K channels. The tail current measurements were obtained in the absence of toxin (open circles) or in the presence of 1 µM Hanatoxin (filled circles).

**Figure 2.** Inhibition of the drk1 K⁺ channel by Hanatoxin. (A) Families of current records obtained from two-electrode voltage-clamp recording of oocytes expressing the wild-type (left) and two mutant drk1 K⁺ channels (F274R, middle; E277K, right), both in the absence and presence of 1 µM toxin. Currents were elicited by voltage steps beginning at -50 mV and incrementing in 5 mV steps from a holding voltage of -80 mV. All traces are shown without subtraction of leak or capacitive currents. (B) Tail current-voltage relations for wild-type (left), F274R (middle), and E277K (right). Tail current amplitudes were measured 2-3 ms after repolarization to -50 mV. All traces are shown without subtraction of leak or capacitive currents. (C) Fraction of uninhibited tail current plotted against test voltage at different toxin concentrations for wild-type, F274R, and E277K. I is the tail current measured in the presence of toxin and I₀ is the tail current measured in the absence of toxin. Data are shown as mean ± SEM for three to six cells at each toxin concentration.
ages (\(I/I_0\)), which approximates the fraction of unbound channels, was used to calculate the \(K_d\) for toxin binding to the \(K^+\) channel (see Materials and methods) (Swartz and MacKinnon, 1997a). In the absence of toxin, the voltage-activation relations for F274R and E277K displayed only small rightward shifts relative to the wild-type channel. In contrast, for both mutants, even very high toxin concentrations produced much less inhibition at negative voltages when compared with the wild-type channel, suggesting that the binding affinity of the toxin for these mutants is greatly diminished.

Fig. 3 shows the results for multiple substitutions at F274, where mutation to Ala was previously reported to reduce Hanatoxin binding affinity by \(\sim 25\)-fold. The 14 substitutions made at F274 include 8 hydrophobic residues (Cys, Ala, Met, Pro, Tyr, Trp, Val, Ile), 2 basic residues (Lys, Arg), 2 acidic residues (Glu, Asp), Ser, and Gly. Fig. 3A shows the dependence of \(I/I_0\) on toxin concentration for the wild-type and two mutant channels, F274G and F274R. For the wild-type channel, the data were well described by a model assuming four equivalent and independent binding sites for Hanatoxin on the drk1 \(K^+\) channel with a \(K_d\) of 103 nM for toxin binding to each site (see materials and methods). The two mutants, F274G and F274R, greatly reduced the toxin binding affinity with \(K_d\) values of 4.4 and 61.6 \(\mu\)M, respectively. The normalized Hanatoxin \(K_d\) values for all 14 substitutions at position 274 are summarized in Fig. 3B. One prominent trend in the data is that substitutions with hydrophobic amino acids cause the smallest perturbation in Hanatoxin binding energy. The range of \(\Delta K_d\) values for hydrophobic residues is 1.2–1.8 kcal mol\(^{-1}\), while the range of \(\Delta K_d\) values for nonhydrophobic residues is 2.4–3.7 kcal mol\(^{-1}\). Although the hydrophobic residues caused the smallest perturbations, even these perturbations were quite significant, implying a very close and specific interaction with the toxin. These results are consistent with an intimate hydrophobic interaction between F274 and the toxin. It is also interesting that at position 274, substitutions with either Lys or Arg produce the largest perturbations, with \(\Delta K_d\) values of 3.7 and 3.6 kcal mol\(^{-1}\), respectively. One possibility is that F274 is positioned close to a basic toxin residue in the channel-toxin complex (see discussion).

We next examined the effects of multiple substitutions at E277. Fig. 4 shows the concentration dependence of \(I/I_0\) for two mutant channels, E277Y and E277K. Compared with the wild-type channels (same data as for Fig. 3), E277Y and E277K displayed greatly reduced binding affinities for Hanatoxin, with \(K_d\) values of 2.5 and 17.3 \(\mu\)M, respectively. The normalized Hanatoxin \(K_d\) values for all eight substitutions at position 277 are summarized in Fig. 4B. An interesting correlation between the magnitude of the effects and the substituted amino acids emerges. Substitutions with basic residues have the largest effects (\(\Delta K_d = 2.8\) kcal mol\(^{-1}\)), while those with neutral residues have more moderate effects (\(\Delta K_d\) from 1.3 to 2.2 kcal mol\(^{-1}\)). Mutation of E277 to Asp produced no discernible change in toxin binding affinity (\(\Delta K_d = 0.04\) kcal mol\(^{-1}\)). The correlation between the charge of the side chain at 277....
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and toxin binding energy is to a first approximation consistent with a through-space electrostatic interaction between this position on the channel and a basic residue on the toxin. However, two observations argue for a more intimate interaction. First, substitutions with various neutral amino acids produce significantly different perturbations, with \(\Delta \Delta G\) values ranging from 1.3 to 2.2 kcal mol\(^{-1}\). Second, the large values of \(\Delta \Delta G\) for all mutations (1.3–2.8 kcal mol\(^{-1}\)), with the exception of Asp, are more consistent with intimate or short-range interactions. We speculate that E277 interacts with a basic residue on Hanatoxin through formation of a salt bridge. In this case, the apparent tranquility observed upon substitution of Asp for Glu would imply some degree of conformational flexibility in either the toxin or the channel.

The results of 13 substitutions at position 273 are shown in Fig. 5. The mutation to Tyr produces the largest change in toxin binding affinity (\(\Delta \Delta G = 2.8\) kcal mol\(^{-1}\)). Substitutions with Thr, Arg, Asp, Ala, Gly, and Cys cause moderate changes (\(\Delta \Delta G = 1.5\)–1.7 kcal mol\(^{-1}\)) in toxin binding affinity, while substitutions with Val, Met, Trp, Phe, His, and Leu produce only minor changes (\(\Delta \Delta G \approx 0.5\) kcal mol\(^{-1}\)). These results are more complex than observed at either F274 or E277. The observation that most hydrophobic substitutions (Val, Met, Trp, Phe, and Leu) produced little change in toxin binding affinity with \(\Delta \Delta G < 0.3\) kcal mol\(^{-1}\) seems to hint at a hydrophobic interaction between position 273 and the toxin. However, the pattern is not simple because mutation to Tyr produces the largest perturbations (\(\Delta \Delta G = 2.8\) kcal mol\(^{-1}\)) and very significant perturbations (1.6 and 1.7 kcal mol\(^{-1}\)) were seen for mutation to Ala and Cys, two smaller hydrophobic residues. A partial explanation might be that I273 is involved in a hydrophobic interaction with the toxin and that there is a requirement for both a large and hydrophobic side-chain at position 273.

Figure 4. Effects of multiple substitutions at position 277 on Hanatoxin binding affinity. (A) Concentration dependence for inhibition of the wild-type and two mutant drk1 K\(^+\) channels by Hanatoxin. \(I/I_0\) is the value of \(I/I_0\) measured in the plateau phase at negative voltages (e.g., Fig. 2 C). Symbols are experimental data for the mean \pm SEM of three to six cells. Solid lines correspond to \(I/I_0 = (1 - \frac{P}{[\text{toxin}]} + K_d)\), with \(K_d\) values of 2.5 and 17.3 \(\mu\)M for E277Y and E277K, respectively. Data for the wild-type channel are the same as in Fig. 3. (B) Normalized \(K_d\) values of eight substitutions at position 277. Mean \pm SEM (n = 3–13) for each mutant channel. The corresponding \(\Delta \Delta G\) values are (kcal mol\(^{-1}\)): 2.8 R, 2.8 K, 2.2 W, 2.0 Y, 1.6 G, 1.5 Q, 1.3 N, and 0.04 D.

The above results are consistent with intimate interactions between three residues in the COOH-terminal end of S3 and Hanatoxin. While the surface area at the toxin-channel interface is unknown, the dimensions of Hanatoxin (~20 \(\times\) 25 \(\AA\)) argue that many channel residues, perhaps 10 or more, are located at the toxin-channel interface (Takahashi et al., 2000). We therefore continued to search for residues contributing to the Hanatoxin receptors by Ala-scanning from the NH\(_2\)-terminal part of S1 through the COOH-terminal Edge of S3 (started at K185 and ended at Y269). Fig. 6 shows the normalized Hanatoxin \(K_d\) values for the 85 mutants made in this region (S1–S3) together with the previous results for an Ala-scan of the S3–S4 linker and the S4 segment (Swartz and MacKinnon, 1997b), shown for comparison. A Kyte-Doolittle hydrophobicity index profile for this region is shown to mark the approximate position of each transmembrane segment (S1–S4). In contrast to the previously observed large perturbations in the COOH-terminal part of S3 (and small changes and in S4), the mutations in S1 through most of S3 caused no more than threefold changes in the toxin \(K_d\). These results suggest that residues from the NH\(_2\)-terminal end of S1 to the NH\(_2\)-terminal part of S3 do not appear to be critical components of the Hanatoxin receptors.
Does Hanatoxin Interact with Residues on the Peripheral Surface of the Pore Domain?

From previous experiments, we know that Agitoxin$_2$, a well-studied pore-blocking toxin, and Hanatoxin can simultaneously occupy the drk1 K$^+$ channel (Swartz and MacKinnon, 1997b). Agitoxin$_2$ belongs to a family of closely related toxins that bind over the central pore axis and have dimensions of $\sim 20 \times 30$ Å (MacKinnon and Miller, 1988; Miller, 1988; Park and Miller, 1992a; Hidalgo and MacKinnon, 1995; Krezel et al., 1995; Ranganathan et al., 1996; MacKinnon et al., 1998). Thus, co-occupancy of the two toxins indicates that the Hanatoxin receptors must be located in peripheral regions on the extracellular surface of the channel, probably at least 15 Å away from the central pore axis (Fig. 7 B). However, from the structure of the KcsA K$^+$ channel (Doyle et al., 1998), we can infer that a large fraction of the extracellular surface of the pore domain lies outside the footprint of Agitoxin$_2$ (MacKinnon et al., 1998) and therefore represents a potential interaction surface for Hanatoxin. To test for a possible interaction between Hanatoxin and the peripheral surface of the pore domain in the drk1 K$^+$ channel, we aligned the sequences for the transmembrane part of KcsA (TM1 through TM2) and the pore domain of the drk1 K$^+$ channel (S5 through S6) (Fig. 7 A). The alignment shows that $\sim 50\%$ of the residues in this region are conserved between the two types of K$^+$ channels. From the alignment, we identified 18 residues on the pore domain of drk1 that correspond to residues located on the perimeter of the extracellular surface of KcsA (Fig. 7 A, asterisks, and B, gray side chains). Most residues were mutated to Ala and the binding affinity of Hanatoxin was determined. As apparent from the normalized Hanatoxin $K_d$ values in Fig. 7 C, the largest change in Hanatoxin binding affinity was slightly less than two-fold. These results argue against an intimate interaction between Hanatoxin and the extracellular surface of the pore domain, and thereby suggest that the toxin binding sites are confined within the voltage-sensing domains of the channel.

DISCUSSION

Previous studies support the idea that Hanatoxin interacts with four receptors on the voltage-gated K$^+$ channel and inhibits the channel by modifying the energetics of gating (Swartz and MacKinnon, 1997a,b). The objective of the present study was to explore two questions. First, what is the molecular composition of the Hanatoxin receptors? Second, where are these receptors located in the structure of the voltage-gated K$^+$ channel?

In the first part of this paper, we examined the effects of multiple substitutions at three positions in the COOH-terminal end of S3, where mutations to Ala were previously found to alter toxin binding affinity. Our results for both I273 and F274 are consistent with hydrophobic interactions between these positions and hydrophobic residues on the toxin. The Glu residue at position 277 also seems to interact intimately with the toxin, perhaps by way of a salt bridge with a basic residue on the toxin. The solution structure of Hanatoxin was recently solved by NMR spectroscopy (Takahashi et al., 2000). The toxin is composed of two $\beta$ strands, one turn of a 3$\alpha$ helix, with four chain reversals. While the surface of the toxin that interacts with the K$^+$ channel
remains to be experimentally identified, a channel interacting face has been proposed based on a structural comparison of different gating modifier toxins (Takahashi et al., 2000). Two types of gating modifier toxins that slow inactivation of Na\(^+\) channels are the α-scorpion toxins and the sea anemone toxins. Hanatoxin and these Na\(^+\) channel toxins each have a face that contains a large hydrophobic patch surrounded by basic and acidic residues. This structural feature is illustrated in the surface rendering of Hanatoxin in Fig. 8 A. Our results would be consistent with F274, and possibly I273, interacting with the hydrophobic cluster on the toxin. E277 might interact with one of the many basic residues (through formation of a salt bridge) that surround the hydrophobic patch. While our results with F274 are consistent with this residue participating in a hydrophobic interaction, they also show that the most dramatic perturbations result from mutations to either Lys or Arg. This might be an indication that in the toxin–channel complex, F274 is located close to a basic toxin residue, a quite feasible scenario for the face of Hanatoxin shown in Fig. 8 A. It is interesting that the face of Hanatoxin proposed to interact with the channel resembles other well-studied protein–protein interfaces. For example, the interface between growth hormone and its receptor shows a patch of hydrophobic residues, where tight hydrophobic contacts exist, surrounded by polar residues that participate in hydrogen bonds and salt bridges (Clackson and Wells, 1995; Bogan and Thorn, 1998; Clackson et al., 1998).

The dimensions of Hanatoxin (\(~20 \times 25\) Å) strongly suggest that many channel residues, certainly more than three, are located at the interaction surface between toxin and channel. To search for additional components of the toxin receptors, we first systematically mutated the region spanning from the NH\(_{2}\) terminus of S1 to the COOH terminus of S4. All mutants in this region displayed binding affinities for Hanatoxin that are comparable to the wild-type channel (Fig. 6). The other region that we examined for potential interactions with Hanatoxin was the extracellular surface of the pore domain. From the known structure of the KcsA K\(^+\) channel, we can infer that the extracellular surface of the pore domain in voltage-gated K\(^+\) channels has sufficient area to accommodate Agitoxin 2, a pore-blocking toxin, and at least portions of four Hanatoxin molecules. When we mutated 18 residues (representing 72 positions in the homotetramer) on the peripheral surface of the pore domain, we found no significant alterations in Hanatoxin binding affinity (Fig. 7). These results argue that Hanatoxin does not interact with the extracellular surface of the pore domain and therefore most likely interacts exclusively with the voltage-sensing domains. Since the extracellular surface of the KcsA K\(^+\)
channel has dimensions of \(~45 \times 45\) Å (Doyle et al., 1996), we conclude that the Hanatoxin receptors must be located at least 20–25 Å away from the central pore axis in voltage-gated K\(^+\) channels, as illustrated by the Hanatoxin footprints in Fig. 8 C.

It is remarkable that mutations of only three residues in the entire voltage-sensing domain (S1–S4) of the drk1 K\(^+\) channel have large effects on Hanatoxin binding affinity. One possibility is that Ala-scanning mutagenesis may have missed residues that contribute either weakly or moderately to the interaction with toxin because some substitutions to Ala represent rather conservative changes. Another very interesting possibility is that the toxin also interacts with the NH\(_2\)-terminal part of S4, where a previous Ala-scan identified several positions where mutations have small effects (three- to five-fold) on Hanatoxin binding affinity (Swartz and MacKinnon, 1997b). We did not examine this region in the present study because the previously reported effects are rather small and therefore difficult to study. However, the large number of “negative” mutations observed elsewhere in the channel strengthens the possibility that there is an important interaction between Hanatoxin and the NH\(_2\)-terminal part of S4. While we probably have not found all the residues that lie at the interface with Hanatoxin, we may have identified the residues (I273, F274, E277) that dominate the energetics of the toxin–channel interaction. This notion is consistent with other examples of protein–protein interactions, where only a few residues can account for most of the binding energy even though many more residues lie at the interface. Two notable examples are the barnase:barnstar and growth hormone:receptor complexes. In both instances, when all residues present at
the protein–protein interfaces are mutated to Ala, only ~25% of them are found to cause substantial perturbations in binding energy (Schreiber and Fersht, 1995; Clackson and Wells, 1995; Bogan and Thorn, 1998; Clackson et al., 1998; Conte et al., 1999).

Several recent studies examining the secondary structure of the first four transmembrane segments (S1–S4) in voltage-gated K⁺ channels suggest that all four segments are membrane-spanning α helices (Monks et al., 1999; Hong and Miller, 2000; Li-Smerin et al., 2000a). The identified interaction of Hanatoxin with the COOH terminus of S3, together with the absence of an interaction with the pore domain, suggests that the toxin interacts with the extracellular surface of this bundle of four helices. However, S1, S2, and the linker between them do not appear to interact strongly with the toxin, suggesting that the Hanatoxin receptors are located over S3 and perhaps S4. The S3 segment is interesting in that only the intracellular or NH₂-terminal two-thirds, up to a conserved proline (P268 in drk1), exhibits α-helical character (Hong and Miller, 2000; Li-Smerin et al., 2000a). While there is no evidence for α-helical structure for the COOH-terminal part of S3, there is evidence for α-helical structure in the nearby linker between S3 and S4 (Li-Smerin et al., 2000a). One possibility is that S3 is composed of two helices separated by a short nonhelical region near P268, as illustrated in Fig. 8 B. In this case, the three residues implicated in forming the Hanatoxin receptors (I273, F274, and E277) would be located either within or near the beginning of the second helix. The spacing of these three residues is compatible with helical structure in this region, with one side of the helix comprising the external surface where intimate interactions with the toxin occur.

It is interesting that gating modifier toxins, perhaps as a general rule, interact with the equivalent region of voltage-gated K⁺, Na⁺, and Ca²⁺ channels. The present results are consistent with an intimate interaction between Hanatoxin and E277 in the drk1 K⁺ channel. Mutations at E1613 in the fourth repeat of the brain IIA Na⁺ channel alter the binding affinity of α-scorpion toxin and sea anemone toxin, two types of toxins that slow inactivation (Rogers et al., 1996). Also, mutation of E1658 in repeat four of the α₁A voltage-gated Ca²⁺ channel disrupts the binding of v-Aga-IVA (Winterfield and Swartz, unpublished observations), a gating modifier toxin for the Ca²⁺ channel (McDonough et al., 1997). In sequence alignments, both E1613 in the Na⁺ channel and E1658 in the Ca²⁺ channel are at positions equivalent to E277 in the drk1 K⁺ channel. This connection between different gating modifier toxins is par-
alleled by several reported examples where gating mod-
ifier toxins interact rather promiscuously with different
types of voltage-gated ion channels. Grammotoxin, a
gating modifier of the α1A voltage-gated Ca2+ channel,
also inhibits the drk1 K+ channel and the binding affin-
ity of grammotoxin to the K+ channel is altered by mu-
tations at 1273, F274, or E277 (Li-Smerin and Swartz,
1998). In addition, Hanatoxin can bind to and inhibit
the α1A Ca2+ channel. Another example is kurtoxin,
an α-scorpion toxin that slows inactivation of the brain
IIA Na+ channel and also inhibits T-type voltage-gated
Ca2+ channels by shifting activation to more depolar-
ized voltages (Chuang et al., 1998). Taken together,
these results suggest that gating modifier toxins inter-
act with a common region of the voltage-sensing do-
main in different voltage-gated ion channels and that
this region probably adopts a well conserved three-
dimensional structure.

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