Structure of an excitatory insect-specific toxin with an analgesic effect on mammals from the scorpion Buthus martensii Karsch

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BmK IT-AP is an excitatory insect-specific \( \beta \)-toxin with analgesic effect from the Chinese scorpion Buthus martensii Karsch (BmK) and consists of 72 residues cross-linked by four disulfide bridges. The crystal structure of BmK IT-AP has been determined at a resolution of 2.6 Å by molecular replacement. Compared with the mammal-selective \( \alpha \)-toxins consisting of 64 residues from the scorpion BmK, the general fold of IT-AP features an additional one-and-a-half turn \( \alpha \)-helix at the C-terminal residues 59–65 with a shifted disulfide bridge Cys38–Cys64. The extension and ‘wiggling’ of the C-terminal segment led to a reshaping of the bioactive surface, including the complete destruction of the active site RC comprising the reverse turn (8–12) and C-terminal residues 58–64, the disappearance of an active surface formed by two aromatic residues Trp38 and Tyr42 and the covering of the conserved aromatic cluster Tyr5, Tyr35 and Trp47, which are all critical for the structure and function of mammal-selective \( \alpha \)-toxins. Bj-xtrIT, the only other excitatory insect-specific toxin whose three-dimensional structure has been determined, is distinct from BmK IT-AP. A functionally important five-residue \( \alpha \)-helix (\( \alpha0 \)) formed by residues 17–21 in Bj-xtrIT is deleted in BmK IT-AP and helix \( \alpha1 \) is immediately connected to Cys16 through two residues Leu17 and Phe18. Accordingly, the functional surface of this region in Bj-xtrIT has also been reshaped in IT-AP, which implies subtle differences between BmK IT-AP and Bj-xtrIT in binding to the receptor, although most other critical residues for structure and function adopt almost identical conformations. The crystal structure of IT-AP also forms a sound basis for further study of the structure–function determinants of the analgesic effect.

1. Introduction

Scorpion toxins acting on voltage-dependent sodium channels in excitable cells are mainly related to human envenomation; they are classified as \( \alpha \)-toxins, which inhibit sodium-current inactivation, and \( \beta \)-toxins, which modify the activation process (Couraud et al., 1982). Scorpion toxins exhibit high selectivity towards animals belonging to different phyla. They have been identified to act on mammals, insects and crustaceans with high specificity (Gordon et al., 1998). The most intriguing property of scorpion toxins is that they can discriminate between insect and mammalian sodium channels. The highly insect-specific toxins are considered to belong to the \( \beta \)-toxins, as they compete with \( \beta \)-toxins for the same receptor-binding site (Gordon et al., 1992). These insect toxins include two distinct subgroups: excitatory and depressant (Zlotkin et al., 1971). Excitatory insect-specific toxins induce a sustained...
contraction paralysis in fly larvae and shift insect Na\(^+\)-channel activation to a more negative membrane potential under voltage-clamp conditions (Zlotkin et al., 1985; Pelhate & Zlotkin, 1982). Depressant insect toxins induce a slow depressant and flaccid paralysis preceded by a short transient phase of contracture. The anti-insect specificity of excitatory toxins has been used to study the properties of the sodium channel (Cestele et al., 1997) in insects and in the development of a new insecticide (Stewart et al., 1991; Gurevitz et al., 1998).

Thus far, a series of excitatory insect-specific toxins have been purified and characterized (Zlotkin et al., 1971; Kopeyan et al., 1990; Ji et al., 1996; Deena et al., 1998; Xiong et al., 1999, Cohen et al., 2004), as shown in Fig. 1. These toxins are 70–76 amino-acid residues long with four highly conserved disulfide bridges, three of which are conserved in both \(\alpha\) - and \(\beta\)-toxins. Although a series of three-dimensional structures of scorpion mammal-specific toxins have been elucidated, only one crystal structure of an excitatory insect-specific toxin, Bj-xtrIT, has been reported to date (Deena et al., 1998), mainly because of the difficulty in obtaining a sufficiently large quantity of highly pure and stable sample. Here, we report the crystal structure of a novel excitatory insect-specific toxin, BmK IT-AP, with an analgesic effect on mammals.

BmK IT-AP (insect toxin analgesic protein) was purified from the venom of the scorpion Buthus martensi Karsch (BmK), which is widely distributed in China and East Asia. BmK IT-AP displays typical excitatory insect-selective toxicity. Interestingly, BmK IT-AP also exhibits an analgesic effect, but is devoid of any toxicity to mice (Xiong et al., 1999). The bioassay showed that its analgesic effect was not parallel to the insect toxicity, which suggested a distinct structure–function determinant for the analgesic effect (Guan et al., 2000). BmK IT-AP consists of 72 amino-acid residues cross-linked by four disulfide bridges and is homologous to all other excitatory insect-specific toxins with sequence identity of 75–82%, except for Bj-xtrIT. BmK IT-AP has a sequence identity of 62% with Bj-xtrIT and differs remarkably in the deletion of a five-residue motif (17–21) after the first cysteine (Cys16) (Fig. 1), which has been identified to be part of the toxin-receptor binding site (Cohen et al., 2004). Thus, BmK IT-AP should be considered as a representative of the subtype of excitatory insect toxins with 72 amino-acid residues. The three-dimensional structure of BmK IT-AP will expand our structural knowledge of the excitatory insect-specific toxins and form a sound structural basis for understanding its dual bioactivity: namely, its toxicity to insects and its analgesic effect on mammals.

Furthermore, a series of crystal structures of toxins from the scorpion BmK, including BmK M1, M2, M4, M7 and M8 (PDB codes 1sn1, 1chz, 1sn4, 1kv0 and 1snb, respectively; Li et al., 1996; He et al., 1999, 2000; Guan et al., 2004), have been determined, all of which belong to the \(\alpha\)-toxins. BmK M1 is highly mammal-selective and has been the subject of extensive analysis by mutagenesis in order to identify the important sites

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**Figure 1**

Sequence alignments of BmK IT-AP with BmK \(\alpha\)-toxins and other scorpion excitatory insect toxins. Conserved secondary-structural elements are shown, of which helix \(\alpha2\) is unique to excitatory insect-specific toxins. Sequences are aligned according to the cysteine frame. The cysteine (highlighted with box in blank background) cross-linked to the last cysteine in the sequence is dramatically shifted from position 12 in \(\alpha\)-toxins to position 38 in BmK IT-AP and in all excitatory insect toxins. AsAHT1 and AsAIII are from the scorpion Androctonus australis Hector, LqQTI1 is from scorpion Leirus quinquemaculatus quinquemaculatus, Lq-IT-1 from the scorpion Buthus judaicus (Possani et al., 1999; Deena et al., 1998; Cohen et al., 2004). BmK M1, M2, M4, M8, IT-AP and IT1 are from the scorpion B. martensi Karsch (He et al., 1999; Li et al., 1996; Luo et al., 1997; Xiong et al., 1999).
for structure and function (Wang et al., 2003; Sun et al., 2002, 2003). The structure of BmK IT-AP as an insect-specific β-toxin in comparison with BmK M1 as a mammal-selective α-toxin will reveal the structural determinants that discriminate these two phylogenetically distinct scorpion toxins.

2. Materials and methods

2.1. Purification and crystallization

BmK IT-AP was purified from the crude venom of BmK as described by Xiong et al. (1999). BmK IT-AP was crystallized at 287 K by the hanging-drop vapour-diffusion method. The hanging drop consisted of 2 µl 20 mg ml⁻¹ BmK IT-AP dissolved in 1 mM acetic acid mixed with 2 µl reservoir solution [1.4 M (NH₄)₂SO₄ and 0.1 M CHES pH 9.5] and 0.6 µl 4 mM Zwittergent 3-10. The drops were left to equilibrate against 600 µl reservoir solution. Within one week, crystals grew to dimensions of 0.3 × 0.15 × 0.15 mm (Guan et al., 2000).

Preliminary crystallographic analysis showed that the crystal belongs to space group P 6422 or P 6222, with unit-cell parameters a = b = 64.97, c = 173.48 Å. There are two molecules in the asymmetric unit and the solvent content is estimated to be 60.5% according to the Matthews coefficient (Vₘ = 3.11 Å³ Da⁻¹; Matthews, 1968).

2.2. Data collection and processing

The diffraction data of BmK IT-AP were collected with wavelength 1.0 Å at room temperature on beamline BL-18B at the Photon Factory, KEK, Tsukuba, Japan using an ADSC Q4R CCD detector. A complete data set to 2.6 Å resolution was integrated, scaled and merged using the HKL package (Otwinowski & Minor, 1997). The final data-collection statistics are listed in Table 1.

2.3. Molecular replacement

The structure of BmK IT-AP was solved by the molecular-replacement method using MOLREP v.7.3 (Vagin & Teplyakov, 1997) from the CCP4 v.4.2 software suite (Collaborative Computational Project, Number 4, 1994). The crystal structure of Bj-xtrIT (PDB code 1bcg) was used as a search model. No atoms were substituted or omitted from the model. Both space group P6₁₂₂ and P6₂₂₂ were used in molecular replacement in the resolution range 29.53–3.0 Å. When P6₂₂₂ was used, the best solution calculated by the rotation function gave a correlation coefficient of 30.0% with an R factor of 56.7% and the subsequent translation function gave a clear solution of two molecules in the asymmetric unit with a correlation coefficient of 43.6% and an R factor of 51.1%. When P6₁₂₂ was used, the rotation and translation function gave a
correlation coefficient of only 22.6% with an $R$ factor of 61.0% and a correlation coefficient of 26.8% with an $R$ factor of 60.0%, respectively. This solution was confirmed by checking the packing using the program O (Jones et al., 1991).

2.4. Model building and refinement

The model was rebuilt using O and refined with the program CNS (Brünger et al., 1998). 10% of the unique reflections were selected randomly to calculate the free $R$ factor (Kleywegt & Brünger, 1996). Following rigid-body refinement, several rounds of simulated-annealing refinement were performed alternating with the calculation of electron-density maps and manual adjustment of the model. Positional and individual $B$ factors were then refined and solvent molecules were identified and added to the model. Some electron densities, which were clearly greater than water molecules in the $2F_o - F_c$ and $F_o - F_c$ maps, were defined as sulfate ions. Finally, energy-minimization refinement with CNS was carried out, followed by restrained refinement with REFMACS (Murshudov et al., 1999) in the final round to further improve the accuracy of the model. The final model, including 1136 protein atoms, 65 water molecules and 11 sulfate ions, was refined to $R_{work} = 20.0\%$ ($R_{free} = 24.9\%$) in the resolution range 29.53–2.6 Å. The final structure was analyzed using PROCHECK (Laskowski et al., 1993).

3. Results and discussion

3.1. Quality of the structure

The two molecules, molecule $A$ and molecule $B$, in the asymmetric unit were well refined and the refinement statistics are given in Table 1. Each molecule contains all 72 amino-acid residues. The Ramachandran plot (Ramachandran & Sasi- sekharian, 1968) reveals that 89.2 and 90.8% of the non-glycine and non-proline residues of molecules $A$ and $B$, respectively, fall in the core region; the remainder are all in the allowed region.

Since the C-terminal stretch 67–72 in molecule $B$ is flexible (Fig. 2), the average $B$ factor of molecule $B$ (34.9 Å$^2$) is significantly higher than that of molecule $A$ (27.2 Å$^2$). Except for the last six residues of molecule $B$, all residues are well defined in the electron-density maps. 11 sulfate ions can be identified in $2F_o - F_c$ and $F_o - F_c$ maps; they mainly contribute to the monomer–monomer interactions and stabilization of the C-terminal segment of molecule $A$ (Figs. 2 and 3).

3.2. Overall structure of BmK IT-AP

The two molecules, $A$ and $B$, in the asymmetric unit are related by a non-crystallographic twofold axis. They interact with each other through three hydrogen bonds, including a direct hydrogen bond ($A10\cdot\cdot\cdotB2\cdot\cdot\cdotNZ$, 3.01 Å) and two sulfate-mediated contacts (Fig. 3). Superposition of molecules $A$ and $B$ shows that except for the C-terminal four residues, their conformations are very similar and the root-mean-square deviation of $C^\alpha$ atoms from residues 1–68 is 0.45 Å (Koradi et al., 1996). The C-terminal stretches 68–72 of these two molecules are in different conformational states: rigid in molecule $A$ but flexible in molecule $B$ (Fig. 3). Since the defined conformations of the C-terminal residues 68–72 are mainly stabilized by five sulfate ions in the crystal, the C-terminal segment of IT-AP should be in a flexible state in solution.

The general fold of IT-AP revealed by the crystal structure can be described as consisting of two parts: the N-sector (residues 1–59) and the C-lobe (residues 60–72) (Fig. 3). The N-sector has the same $\beta\alpha\beta\beta$ motif as in $\alpha$-toxins (Fig. 4). It is composed of a dense core of secondary-structure elements, including an $\alpha$-helix formed by residues 19–28 and a three-stranded antiparallel $\beta$-sheet formed by residues 2–5, 32–38 and 45–54. Three disulfide bridges formed by Cys16–Cys37, Cys22–Cys42 and Cys26–Cys44 stabilize this special motif; they are strictly conserved in all scorpion $\alpha$-toxins and $\beta$-toxins. Compared with the $\alpha$-toxins, the only structural change in this part is that loop 38–44, which was identified to be of functional importance in $\alpha$-toxins.
(Sun et al., 2003), is deleted in the sequence of IT-AP (Figs. 1 and 4). However, the C-lobe (residues 60–72) of IT-AP is strikingly different from that in α-toxins; it features an additional one-and-a-half turn α-helix (α2) formed by residues 59–65 and a disulfide bridge Cys38–Cys64 that is shifted from the fourth disulfide bond Cys12–Cys63 of α-toxins (Fig. 4). These new structural elements give IT-AP a molecular framework and topology distinct from that of α-toxins.

3.3. Main structural disparity between BmK IT-AP and BmK α-toxins

In all α-toxins a five-residue reverse turn (8–12) at the N-terminus connects with the C-terminal segment 58–64 to form a unique tertiary structure (Fig. 4), which is anchored to the βαββ molecular scaffold through a complicated hydrogen-bonding network and the fourth disulfide bond Cys12–Cys63.

This unique local structure has been called site RC (He et al., 1999; Wang et al., 2003). Site-directed mutagenesis investigations of BmK M1 have identified site RC as being crucial for the structure and function of the toxin (Wang et al., 2003; Sun et al., 2002). Residues Asn11, Arg58 and Cys12 and Cys63 of the disulfide bridge have been shown to be responsible for stabilizing the unique conformation of this functional site. Mutating any one of them seriously reduced the bioactivity of the toxin and synchronously altered the conformation (Wang et al., 2003; Sun et al., 2002). In fact, in all α-toxins Asn11 provides side-chain atoms O^1 and N^2 to connect the main chain of Val59 through 11 O^1–11 N^2 hydrogen bonds (Fig. 4). In addition, the carbonyl O atom of Asn11 makes a hydrogen bond with Arg58 N$^\prime$, which is further connected to Gly61 via a hydrogen bond. Correspondingly, Asn11 and Cys12 are strictly conserved and residue 58 is conserved as Arg or Lys in all α-toxins (Cestele et al., 1999). Furthermore, mutagenesis analysis (Sun et al., 2002) has shown that the disulfide bond Cys12–Cys63 is only critical for stabilizing the local structure relevant to the C-tail in BmK M1, not for the general folding of M1. In the structure of IT-AP, residue 11 is changed to Gly from Asn (Fig. 1) and all the interactions described above in the α-toxins are lost. Additionally, the fourth disulfide bridge is shifted from 12–63 to a new position 38–64 to connect the lengthened C-terminal segment to the molecular scaffold with a new pattern (Fig. 4). It seems that the change of residue 11 from Asn to Gly is a critical structural factor for switching on the unique C-tail conformation of BmK IT-AP.

Site-directed mutagenesis of BmK M1 also suggested that α-toxins have two bioactive surfaces which may directly interact with the receptor site on the sodium channel. One is located in site RC and mainly consists of the C-terminal basic residues Lys62 and His64 (Wang et al., 2003). Another one involves two conserved aromatic residues Trp38 and Tyr42 on the loop 38–44 between β2 and β3, called site AR (Sun et al., 2003). In BmK IT-AP the first active surface has clearly been changed. The hydrogen-bonding network between the turn 8–12 and the C-terminus in site RC of BmK M1 has been destroyed completely in IT-AP (Fig. 4), so the surface of IT-AP in this region differs distinctly from that of M1 (Fig. 5). Furthermore, in IT-AP the residues corresponding to residues

![Figure 4](image-url)

Comparison of BmK IT-AP and BmK α-toxins (α-like toxin M1 and classic α-toxin M8). (a) Overall structures of BmK IT-AP, M1 (He et al., 1999) and M8 (Li et al., 1996). C-termini are shown in magenta, where IT-AP forms a unique helix distinct from those of M1 and M8. Three conserved disulfide bridges and the shifted disulfide bridge (S4) are highlighted in yellow. The conserved common structure motif βαββ is shown in red and green. (b) The local structures of the reverse turn 8–12 (RT) and the C-terminal segment (CT) in BmK M1 and in IT-AP, showing some of the structural factors that trigger the unique C-terminal conformation of BmK IT-AP. In M1 the turn 8–12 connects the C-terminus 58–64 via a hydrogen-bonding network in which Asn11 plays a critical role to form a unique tertiary arrangement (site RC). In IT-AP the network is absent; Asn11 in M1 is mutated to Gly11 in IT-AP, the disulfide bridge Cys12–Cys63 in M1 is shifted to Cys38–Cys64 and the orientation of the C-terminus is dramatically changed to be distant from the turn 8–12.
41–45 of the α-toxins have been deleted and Trp38 has changed to Leu, which leads to the disappearance of another active surface site AR in α-toxins (Figs. 4 and 5).

Furthermore, in all α-toxins there is a conserved aromatic residue cluster, Tyr5, Tyr35 and Trp47, apparent on the so-called the conserved hydrophobic surface (CHS), which is assumed to be part of the functional site targeting the sodium channel (Fontecilla-Camps et al., 1988; Li et al., 1996). Detailed mutagenesis analysis of these three aromatic residues has shown that they contribute to the function of the toxin mainly through stabilizing the three-stranded β-sheet (Sun et al., 2003). In the IT-AP structure, this aromatic cluster still exists, but it is covered by the lengthened C-terminal segment (Fig. 5). This indicates that this conserved aromatic residue cluster, while still able to stabilize the molecular scaffold in BmK IT-AP, is not involved in interaction with the receptor.

Evidently, although the α-toxins and β-toxins possess the same molecular scaffold in the N-terminal part, the extension and ‘wiggling’ of the C-terminal segment of BmK IT-AP have led to reshaping of the bioactive surface and a different topology of the molecule; this forms the main structural basis for the insect-specific bioactivity of BmK IT-AP. It is plausible to infer that the functional site of IT-AP will be completely different from that of α-toxins.

### 3.4. Comparison of BmK IT-AP with Bj-xtrIT

The only other crystal structure of an excitatory insect-specific toxin determined to date is that of Bj-xtrIT from the scorpion Butholus judaicus (Deena et al., 1998). Bj-xtrIT shares 44–62% sequence identity with other excitatory toxins found so far, including BmK IT-AP, and differs remarkably from the others, with an insertion of five residues (17–21) after the first conserved cysteine (Cys16). The superimposed main chains of BmK IT-AP and Bj-xtrIT (Fig. 6) show that the N-terminal motif and the additional helix α2 in the C-terminal part are similar in the two toxins. The major differences between BmK IT-AP and Bj-xtrIT occur at the five-residue insertion (17–21, numbering as in Bj-xtrIT) (Fig. 1) after the first conserved cysteine (Cys16) and prior to the first helix α1. The crystal structure of Bj-xtrIT showed that these inserted residues formed a short α-helix (α0) on the molecular surface. More recently, site-directed mutagenesis has identified the solvent-exposed region formed by these residues as part of the surface of Bj-xtrIT that interacts with its receptor site (Cohen et al., 2004). In the BmK IT-AP structure, this short helix (α0) is absent and helix α1 is directly connected to Cys16 through two residues Leu17 and Phe18 (numbering as in BmK IT-AP). Accordingly, the functional surface in this region of Bj-xtrIT has also been reshaped in IT-AP (Fig. 6). Therefore, it is plausible to infer that the property of binding to the receptor site on the sodium channels may be subtly different between BmK IT-AP and Bj-xtrIT. Therefore, the structures of BmK IT-AP and Bj-xtrIT could be considered as representative research papers.
models of two subtypes of the excitatory insect-specific toxins.

Dissection of the functional sites of Bj-xtrIT by mutagenesis has also suggested that two residue clusters constitute the main functional surface of the toxin (Cohen et al., 2004). One cluster appears in helix $\alpha 1$ and its vicinity, including Glu15, Glu30, Tyr26 and Val34, and the other cluster in the C-terminal region, including Val71, Gln72, Ile73 and Ile74 (Fig. 6). These residues invariably appear at the corresponding positions of BmK IT-AP. The structural comparison indicates that residues Glu15, Glu30, Tyr26 and Val34 adopt almost identical conformations in both IT-AP and Bj-xtrIT (Fig. 6). The residue cluster at the C-terminal region showed some conformational difference between the two toxins. However, the C-terminal segments also displayed distinct conformational states for molecules A and B in the asymmetric unit owing to different environments in the crystal, as described above. It seems that the flexibility of the C-terminal stretch is intrinsic. Thus, the conformational difference of the C-terminal segments observed in BmK IT-AP and Bj-xtrIT may not have biological significance.

### 3.5. Analgesic effect of BmK IT-AP

Two excitatory insect toxins, IT-AP and AngP1 (Xiong et al., 1999; Guan et al., 2001), from the venom of the scorpion BmK have been found to produce an analgesic effect on mice. However, the mechanism by which these toxins modulate the sensation of pain remains to be clarified. Evidently, the analgesic effect of IT-AP and AngP1 cannot be ascribed to any mammalian neurotoxicity the peptides may possess, as they have no detectable toxicity to mice. In fact, the present structure of IT-AP showed that all the functional sites of the BmK toxins specific to mammals have been changed in IT-AP, as described above. Additionally, a bioassay indicated that the analgesic effect of IT-AP on mice is at least 4–5 times stronger than that of AngP1, but its toxicity to insects is two times weaker than that of AngP1 (Guan et al., 2001). Evidently, the analgesic effects of these peptides are not parallel to the insect toxicity. Thus, it seems that IT-AP and AngP1 may have unique structure–function determinants responsible for the analgesic effect that are distinct from those for the insect-specific effects. The crystal structure of BmK IT-AP forms a sound basis for further study on the structural mechanism of the analgesic effects of IT-AP.

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### References


