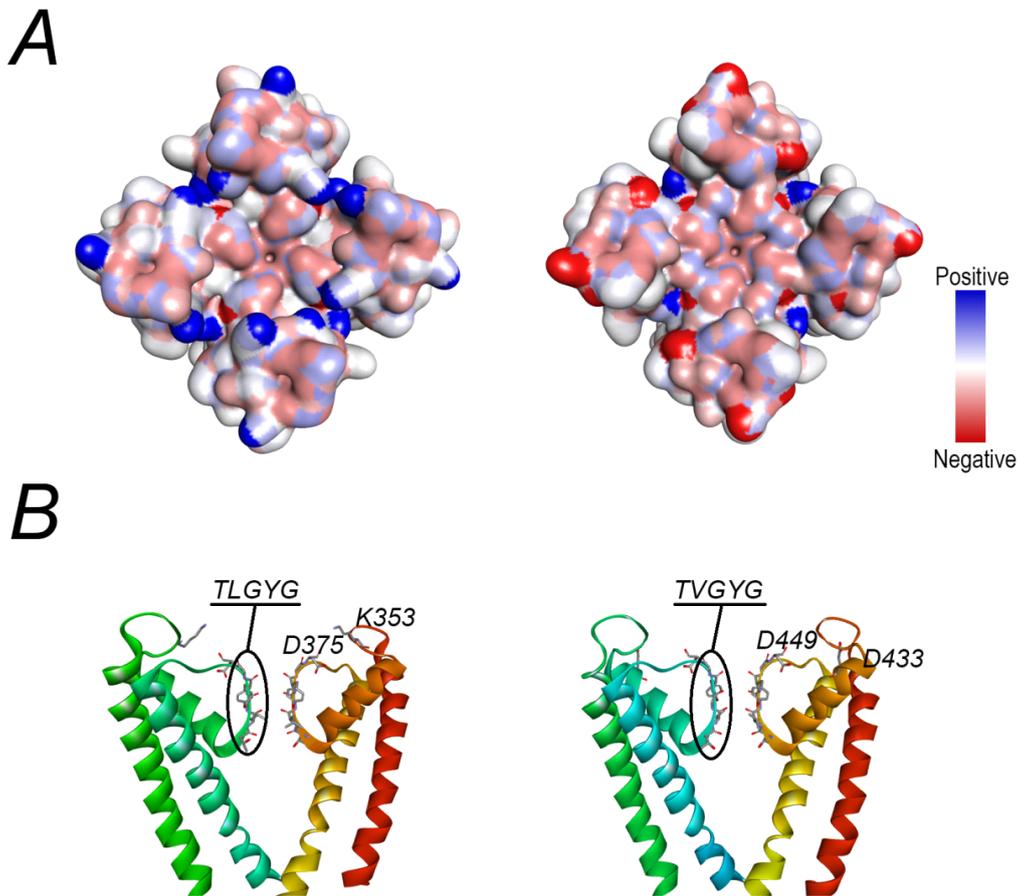
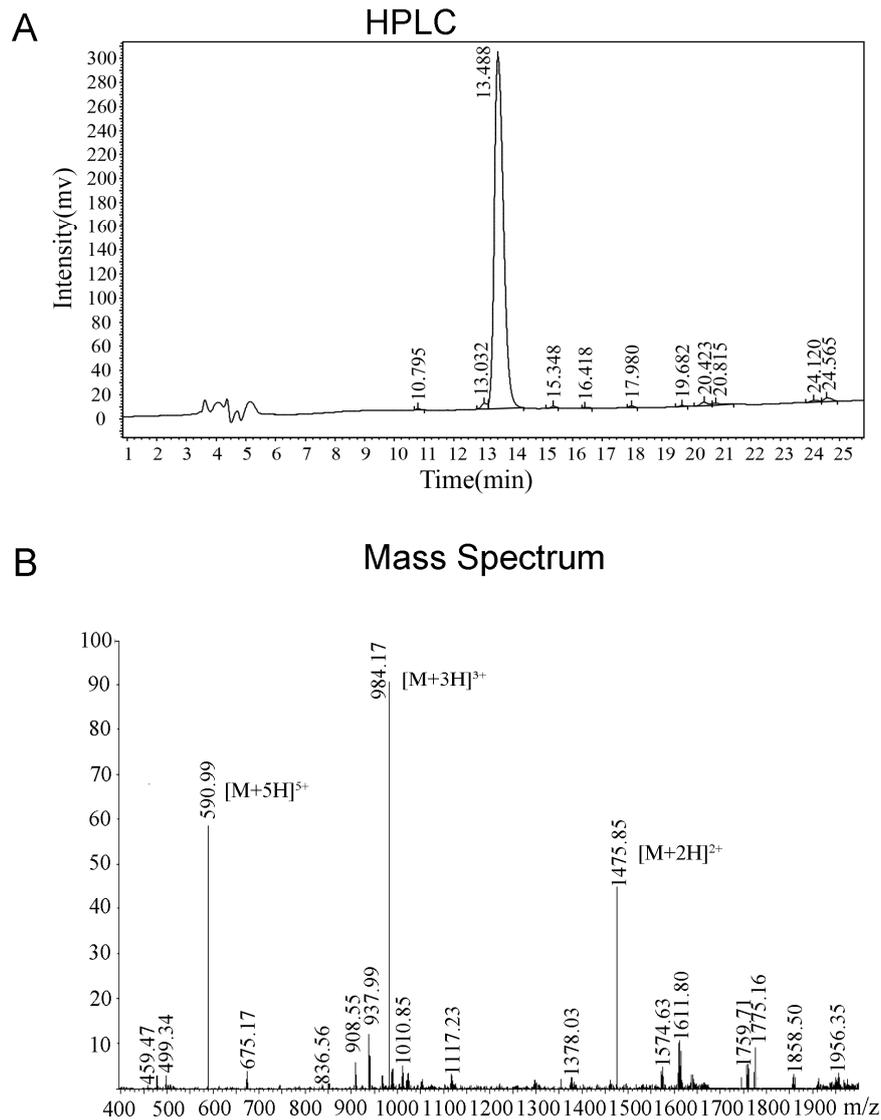


# Supplementary Materials: BmP02 Atypically Delays Kv4.2 Inactivation: Implication for a Unique Interaction Between Scorpion Toxin and Potassium Channel

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**Figure S1.** Homology modeling of Kv4.2 and Kv1.3 using the structure of KcsA (PDB: 1BL8) as template. **(A)** A vertical view of Kv4.2 (left) and Kv1.3 (right). The positively charged groups are shown in blue and the negatively charged groups are in red; **(B)** An insight into the structure of Kv4.2 (left) and Kv1.3 (right). Residues in the selectivity filter and a Asp nearby are highlighted. K<sub>353</sub> in Kv4.2 and D433 (the corresponding residue of A<sub>359</sub> in Kv4.2) in Kv1.3 are also highlighted.



**Figure S2.** The purity and the molecular weights of synthetic BmP02. **(A)** Purity of synthetic BmP02 was determined by High Performance Liquid Chromatography (HPLC); **(B)** Molecular weights of the peptides were determined by mass spectrum (MS).

**Table S1.** The primers used in the construction of mutants. The mutated sites are underlined. S: sense, A: anti-sense.

Name	Mutated Sites	Primer
Kv4.2M1	A <sub>359</sub> D	S:5'-CAGCATCCCTGACGCCTTCTGGTATAACCATCGT A:5'-GGTATACCAGAAGGCGTCAGGGATGCTGGTGAA
Kv4.2M2	K <sub>347</sub> A/K <sub>353</sub> G	S:5'-GCGGGGTCTTCGGCTAGCGGGTTCACCAGCATCCCT A:5'-CCCGTAGCCGAAGACCCCGCTCTGCGTAGAACAT
Kv4.2M3	K <sub>347</sub> A/K <sub>353</sub> G/A <sub>359</sub> D	S:5'-GCGGGGTCTTCGGCTAGCGGGTTCACCAGCATCCCTGACGCCTTCTGGTA TACCATCGT A:5'-GTCAGGGATGCTGGTGAACCCGCTAGCCGAAGACCCCGCTCTGCGTAG AACATAACTG