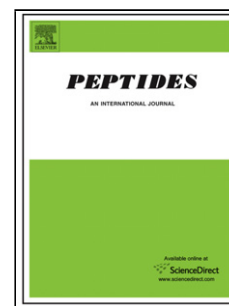


## Accepted Manuscript

Title: Structure, folding and stability of a minimal homologue from *Anemonia sulcata* of the sea anemone potassium channel blocker ShK

Authors: Bankala Krishnarjuna, Christopher A. MacRaild, Punnepalli Sunanda, Rodrigo A.V. Morales, Steve Peigneur, Jason Macrander, Heidi H. Yu, Marymegan Daly, Srinivasarao Raghothama, Vikas Dhawan, Satendra Chauhan, Jan Tytgat, Michael W. Pennington, Raymond S. Norton



PII: S0196-9781(17)30305-4  
DOI: <https://doi.org/10.1016/j.peptides.2017.10.001>  
Reference: PEP 69838

To appear in: *Peptides*

Received date: 7-7-2017  
Revised date: 3-10-2017  
Accepted date: 5-10-2017

Please cite this article as: Krishnarjuna Bankala, MacRaild Christopher A, Sunanda Punnepalli, Morales Rodrigo AV, Peigneur Steve, Macrander Jason, Yu Heidi H, Daly Marymegan, Raghothama Srinivasarao, Dhawan Vikas, Chauhan Satendra, Tytgat Jan, Pennington Michael W, Norton Raymond S. Structure, folding and stability of a minimal homologue from *Anemonia sulcata* of the sea anemone potassium channel blocker ShK. *Peptides* <https://doi.org/10.1016/j.peptides.2017.10.001>

This is a PDF file of an unedited manuscript that has been accepted for publication. As a service to our customers we are providing this early version of the manuscript. The manuscript will undergo copyediting, typesetting, and review of the resulting proof before it is published in its final form. Please note that during the production process errors may be discovered which could affect the content, and all legal disclaimers that apply to the journal pertain.

## Structure, folding and stability of a minimal homologue from *Anemonia sulcata* of the sea anemone potassium channel blocker ShK

Bankala Krishnarjuna,<sup>a</sup> Christopher A. MacRaid,<sup>a</sup> Punnepalli Sunanda,<sup>b</sup> Rodrigo A. V. Morales,<sup>a</sup> Steve Peigneur,<sup>c</sup> Jason Macrander,<sup>d,e</sup> Heidi H. Yu,<sup>f</sup> Marymegan Daly,<sup>d</sup> Srinivasarao Raghothama,<sup>b</sup> Vikas Dhawan,<sup>g</sup> Satendra Chauhan,<sup>f</sup> Jan Tytgat,<sup>c</sup> Michael W. Pennington,<sup>g</sup> and Raymond S Norton,<sup>a\*</sup>

<sup>a</sup> Medicinal Chemistry, Monash Institute of Pharmaceutical Sciences, Monash University, 381 Royal Parade, Parkville, VIC 3052, Australia.

<sup>b</sup> NMR Research Centre, Indian Institute of Science, Bangalore 560012, India.

<sup>c</sup> Toxicology and Pharmacology, University of Leuven, O&N 2, Herestraat 49, P.O. Box 922, 3000, Leuven, Belgium.

<sup>d</sup> Department of Evolution, Ecology, and Organismal Biology, Ohio State University, 1315 Kinnear Rd, Columbus, OH 43212, USA.

<sup>e</sup> Department of Biology, University of North Carolina at Charlotte, 9201 University City Blvd, Charlotte, NC 28223

<sup>f</sup> Infection and Immunity Program, Monash Biomedicine Discovery Institute and Department of Microbiology, Monash University, VIC 3800, Australia.

<sup>g</sup> Peptides International, Louisville, Kentucky 40299, USA.

\*Correspondence to: Professor Raymond S. Norton, Medicinal Chemistry, Monash Institute of Pharmaceutical Sciences, Monash University, 381 Royal Parade, Parkville, Victoria 3052, Australia. Phone: (+61 3) 9903 9167. Fax: (+61 3) 9903 9582 E-mail: ray.norton@monash.edu

### Graphical abstract

AsK132958 is a 29-residue peptide identified in a transcriptomic study of *Anemonia sulcata*. It has the same disulfide framework and a similar structure to ShK. AsK132958 is not active against Kv1.3 channels, owing to the lack of a Lys-Tyr dyad and other functionally important amino acid residues. AsK132958 is more resistant to proteolysis than ShK. Introducing a Lys-Tyr functional dyad to the AsK132958 structural scaffold may be a useful way of developing a proteolytically stable Kv1.3 blocker.

Download English Version:

<https://daneshyari.com/en/article/8347609>

Download Persian Version:

<https://daneshyari.com/article/8347609>

[Daneshyari.com](https://daneshyari.com)