

## VSTx-3

<b>Cat. No.:</b>	HY-P5916
<b>Molecular Formula:</b>	C <sub>182</sub> H <sub>261</sub> N <sub>51</sub> O <sub>51</sub> S <sub>6</sub>
<b>Molecular Weight:</b>	4171.72
<b>Sequence:</b>	Asp-Cys-Leu-Gly-Trp-Phe-Lys-Gly-Cys-Asp-Pro-Asp-Asn-Asp-Lys-Cys-Cys-Glu-Gly-Tyr-Lys-Cys-Asn-Arg-Arg-Asp-Lys-Trp-Cys-Lys-Tyr-Lys-Leu-Trp (Disulfide bridge: Cys2-Cys17, Cys9-Cys22, Cys16-Cys29)
<b>Sequence Shortening:</b>	DCLGWFKGCDPDNDKCCEGYKCNRRDKWCKYKWLW (Disulfide bridge: Cys2-Cys17, Cys9-Cys22, Cys16-Cys29)
<b>Target:</b>	Others
<b>Pathway:</b>	Others
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.

## BIOLOGICAL ACTIVITY

### Description

VSTx-3 is a K<sub>V</sub> channel blocker. VSTx-3 is demonstrated to be a potent, TTX-sensitive sodium channel blocker and especially, a potent blocker of NaV1.8 channels (IC<sub>50</sub> 0.19 μM for hNaV1.3, 0.43 μM for hNaV1.7 and 0.77 μM for hNaV1.8 channels).

## REFERENCES

[1]. Ronit S Cherkj, et al. Two tarantula venom peptides as potent and differential Na(V) channels blockers. *Toxicon*. 2014, 58.

**Caution: Product has not been fully validated for medical applications. For research use only.**

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: [tech@MedChemExpress.com](mailto:tech@MedChemExpress.com)

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA