

## Hm1a

<b>Cat. No.:</b>	HY-P5183
<b>Molecular Formula:</b>	C <sub>170</sub> H <sub>239</sub> N <sub>47</sub> O <sub>54</sub> S <sub>6</sub>
<b>Molecular Weight:</b>	3997.39
<b>Sequence:</b>	Glu-Cys-Arg-Tyr-Leu-Phe-Gly-Gly-Cys-Ser-Ser-Thr-Ser-Asp-Cys-Cys-Lys-His-Leu-Ser-Cys-Arg-Ser-Asp-Trp-Lys-Tyr-Cys-Ala-Trp-Asp-Gly-Thr-Phe-Ser (Disulfide bridge: Cys2-Cys16; Cys9-Cys21; Cys15-Cys28)
<b>Sequence Shortening:</b>	ECRYLFGGCSSTSDCCKHLSCRSDWKYCAWDGTFS (Disulfide bridge: Cys2-Cys16; Cys9-Cys21; Cys15-Cys28)
<b>Target:</b>	Sodium Channel
<b>Pathway:</b>	Membrane Transporter/Ion Channel
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.

### BIOLOGICAL ACTIVITY

#### Description

Hm1a is a disulfide-rich spider-venom peptide, and a NaV1.1 activator. Hm1a restores the function of inhibitory interneurons in Dravet syndrome (DS) mouse model<sup>[1]</sup>.

### REFERENCES

[1]. Chow CY, et al. A selective NaV1.1 activator with potential for treatment of Dravet syndrome epilepsy. *Biochem Pharmacol.* 2020 Nov;181:113991.

**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

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