



ShK

Product name : ShK	Synonyms:
Catalog#: 08SHK001	

Product description

ShK (Stichodactyla helianthus Neurotoxin) has been isolated from the venom of the Carribean sea anemone Stoichactis helianthus. **ShK** inhibits voltage-dependent potassium channels. It blocks $K_v1.3$ (KCNA3) potently and also $K_v1.1$ (KCNA1), $K_v1.4$ (KCNA4), and $K_v1.6$ (KCNA6) respectively with 11 pM,16 pM and 165 pM. Interestingly, it was also demonstrated that **ShK potently inhibits the hK_v3.2b channel** with an IC₅₀ value of approximately 0.6 nM.

Product specifications

AA sequence: Arg-Ser-Cys³-lle-Asp-Thr-lle-Pro-Lys-Ser-Arg-Cys¹²-Thr-Ala-Phe-Gln-Cys¹⁷-Lys-His-Ser-Met-Lys-Tyr-Arg-

Leu-Ser-Phe-Cys²⁸-Arg-Lys-Thr-Cys³²-Gly-Thr-Cys³⁵-OH **Disulfide bonds:** Cys³-Cys¹⁵, Cys¹²-Cys³⁸, Cys¹⁷-Cys³²

Length (aa): 35

Formula: C₁₆₉H₂₇₄N₅₄O₄₈S₇ Appearance: White lyophilized solid Molecular Weight: 4051.87 Da CAS number: 165168-50-3

Source: Synthetic **Counterion:** TFA salts

Solubility: Water or saline buffer, 5 mg/mL maximum (recommendation)

Formulation

Storage/Stability: Shipped at ambient temperature under lyophilized powder. Store at -20°C (-4°F). Do not freeze-thaw. Aliquot sample if required and store at -80°C (-112°F).

Expiry date: One year

Use restrictions: For laboratory use only. Not for drug, household or other uses. Not for use in diagnostic or therapeutic procedures.

Related products

- Kaliotoxin 1 #08KTX002: potent inhibitor of K_v1.1, K_v1.2, K_v1.3 channels
- Margatoxin #08MAG001: selective inhibitor of K_v1.3
- HsTx1 # 08NEU001: inhibits K_v1.3 with a Kd close to 10pM
- <u>Maurotoxin #08MAR001:</u> inhibits voltage-gated potassium channels and small conductance calcium-activated channels
- ADWX-1 #13ADW001: selective K_v1.3 blocker
- (Dap²²)-ShK #13SHD001: selective blocker of the voltage-gated potassium channel K_v1.3 (IC₅₀ ~ 23 pM)

References

- Panyi G, et al. (2006) K+ channel blockers: novel tools to inhibit T cell activation leading to specific immunosuppression. Curr Pharm Des.
- Yan L., et al. (2005) Stichodactyla helianthus peptide, a pharmacological tool for studying Kv3.2 channels. Mol Pharmacol.
- Beeton C., et al. (2005) Targeting effector memory T cells with a selective peptide inhibitor of Kv1.3 channels for therapy of autoimmune diseases. Mol Pharmacol.
- Norton RS., et al. (2004) Potassium channel blockade by the sea anemone toxin ShK for the treatment of multiple sclerosis and other autoimmune diseases. Curr Med Chem
- Castaneda, O., et al. (1995) Characterization of a potassium channel toxin from the Caribbean Sea anemone Stichodactyla helianthus, Toxicon.

For laboratory research use only