



Molecular Tools for the Life Science Community

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Certificate of Analysis

PRODUCT # RTS-370
 LOT # ST-100

rScyllatoxin (Leiurotoxin I, LeTx I)

(*Leiurus quinquestratus hebraeus*)

M.W.: 3424 daltons.
Sequence: AFCNLRMCQL SCRSLLGLLK CIGDKCECVK H
Accession #: [P16341](#)
Purity: > 98%, by HPLC.
Solubility: Any aqueous buffer.

Preparation:

rScyllatoxin is a recombinant peptide expressed in and extracted from *E. coli* and purified to homogeneity.

Reconstitution:

The peptide concentration and identification were determined by amino acid analysis. Each vial contains 5 µg, 10 µg or 0.1 mg of unbuffered protein. Dissolving of 10 µg in 2.92ml of any conventional buffer gives a stock solution of 1 µM. Before dissolving the toxin, the tube should first be centrifuged, to concentrate the lyophilized toxin in the bottom of the tube. After centrifuging, the toxin must be dissolved into a stock solution using distilled water, or an appropriate buffer, to a concentration of 10⁻⁵-10⁻⁶M.

Storage and Stability:

Lyophilized form: 2-3 weeks at room temperature.
 One year at -20° C.
 Liquid form: Up to two weeks at 4° C.
 Three months at -20° C.

Known action:

Scyllatoxin is a 31 amino acid long toxin, originally isolated from the venom of the scorpion *Leiurus quinquestratus hebraeus*, and is classified as α-KTx 5.1 scorpion toxin family, having three disulfide bridges.^{1,2}

Scyllatoxin was shown to compete with [¹²⁵I]-apamin binding in the brain.³ Furthermore, Scyllatoxin appears to be selective for apamin-sensitive SK channels. Scyllatoxin inhibits apamin-sensitive SK channel activity in guinea-pig and rabbit hepatocytes,⁴ SK currents in human lymphoblastoma cells,^{5,6} and epinephrine-induced relaxation of visceral smooth muscle.⁷ Scyllatoxin also inhibits the apamin-sensitive afterhyperpolarization that follow action potentials in skeletal muscle⁷ and neurons.⁸ The SK channel-mediated Afterhyperpolarising current (I_{AHP}) of dorsal vagal neurons, presuming K_{Ca}2.3 (SK3), were blocked by Scyllatoxin (20—30nM).⁹ HEK 293 cell currents stably expressing hK_{Ca}2.1 (hSK1) and K_{Ca}2.2 (hSK2) were blocked by scyllatoxin with IC₅₀ of 80 nM and 287 pM, respectively.¹⁰

Scyllatoxin seems to be a very potent blocker of either K_{Ca} channels with highest affinity towards K_{Ca}2.2, having the following affinities: K_{Ca}2.2 < K_{Ca}2.3 < K_{Ca}2.1.

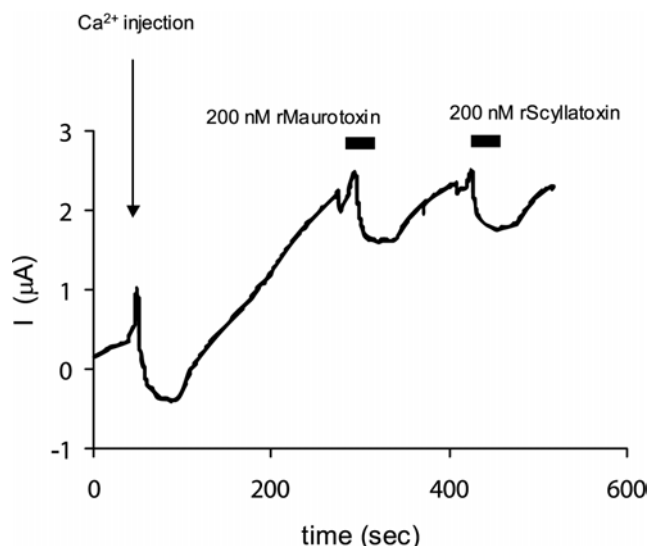


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Bioassay: Inhibition of $K_{Ca2.2}$ channels expressed in *Xenopus* oocytes by rScyllatoxin and by rMaurotoxin.

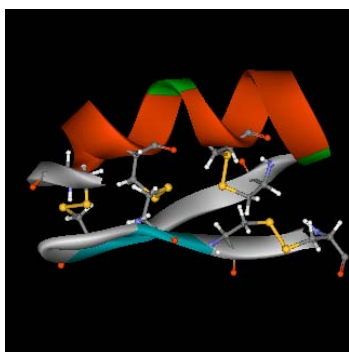


Legend:

Inhibition of $K_{Ca2.2}$ channels expressed in *Xenopus* oocytes by rScyllatoxin and by rMaurotoxin. Continuous current recording at holding potential +5 mV with low Cl^- content in the bath solution. An outward current (upward deflection) carried out by K^+ ions flowing via $K_{Ca2.2}$ channels develops following an intra-oocyte Ca^{2+} injection (arrow). Both recombinant toxins partially and reversibly depressed the K^+ current at 200 nM. Periods of toxin perfusion are marked by the horizontal bars.

References:

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NMR structure of scyllatoxin (1SCY).¹¹ Scyllatoxin belongs to the α -KTX-5.2 scorpion toxin family.²