

Labeled α -Bungarotoxin

α -Bungarotoxin is a 74 amino acid peptidyl toxin isolated from the venom of the banded krait snake, *Bungarus multicinctus*¹.

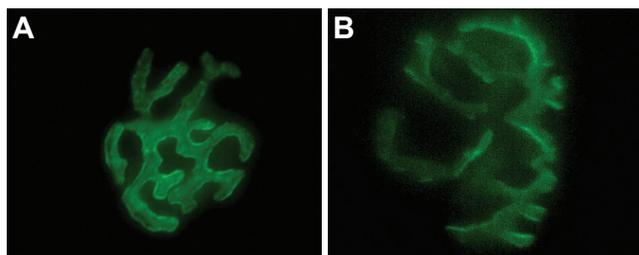
α -Bungarotoxin blocks postsynaptic neuromuscular transmission via competitive inhibition of nicotinic ACh receptors (nAChRs) with an IC_{50} of 3.5×10^{-10} M, thereby prevents the depolarizing action on postsynaptic membranes and blocks neuromuscular transmission².

The toxin is selective for nAChR $\alpha 7$ receptors (IC_{50} value of 1.6 nM) and nAChR $\alpha 3/\beta 4$ receptors (IC_{50} value of $> 3 \mu M$)^{3,4}. α -Bungarotoxin also binds to and blocks a subset of GABA_A receptors (GABA_ARs) that contain the GABA_AR $\beta 3$ subunit. In particular, α -Bungarotoxin blocks GABA_ARs that contain interfaces between adjacent $\beta 3$ subunits⁵.

Labeled α -Bungarotoxin is ideally used to label nAChR $\alpha 7$ receptors, and in many cases to localize neuromuscular junctions^{6,7}.

Alomone Labs is pleased to offer **α -Bungarotoxin-ATTO-488** (#B-100-AG), labeled with a green fluorescent dye and **α -Bungarotoxin-Biotin** (#B-100-B), labeled with biotin.

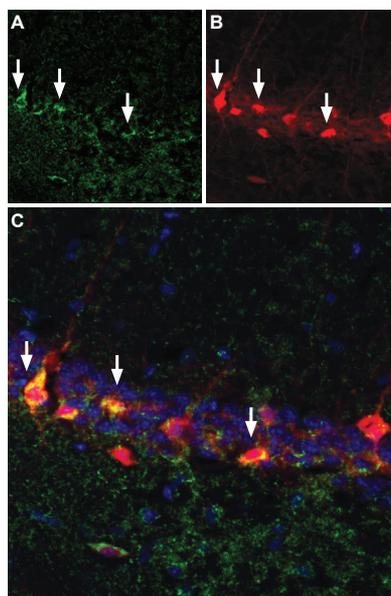
Alomone Labs α -Bungarotoxin-ATTO-488 in whole mount staining of mice Gastrocnemius muscle



Whole mount staining of mice Gastrocnemius muscle was stained with the neuromuscular junction marker **α -Bungarotoxin-ATTO-488** (#B-100-AG), (green) at 1:50 (A) and 1:100 (B) ratios.

The images were taken using Nikon Epifluorescence microscopy at X100 magnification and are a kind gift from Dr. Eran Perlsson, Dept. of Physiology and Pharmacology, Tel-Aviv University.

Bungarotoxin-Biotin binding sites co-localize GABAergic neurons expressing parvalbumin in mouse hippocampal CA1 region



A. Free floating mouse brain sections were incubated with **α -Bungarotoxin-Biotin** (#B-100-B), (1:10,000) followed by streptavidin-Alexa 488 (green). B. Same sections were stained with anti-parvalbumin, followed by goat anti-mouse labeled with Texas red isothiocyanate (TRITC). C. Merge of A and B demonstrates sites of colocalization (vertical arrows) confirming reports that a subset of hippocampal interneurons express nAChR $\alpha 7$. DAPI is used as the counterstain.

References

- Ohta, M. *et al.* (1987) *FEBS Lett.* **222**, 79.
- Wilson, P.T. *et al.* (1988) *Mol. Pharmacol.* **34**, 643.
- Wilson, S.P. and Kirshner, N. (1977) *J. Neurochem.* **28**, 687.
- Garcia-Guzman, M. *et al.* (1995) *Eur. J. Neurosci.* **7**, 647.
- McCann, C.M. *et al.* (2006) *Proc. Natl. Acad. Sci. U.S.A.* **103**, 5149.
- Fertuck, H.C. and Salpeter, M.M. (1974) *Proc. Natl. Acad. Sci. U.S.A.* **71**, 1376.
- Ravdin, P. and Axelrod, D. (1977) *Anal. Biochem.* **80**, 585.

Related Products

Compound	Cat. #
α -Bungarotoxin	B-100
α -Bungarotoxin-ATTO-488	B-100-AG
α -Bungarotoxin-Biotin	B-100-B
Anti-Nicotinic Acetylcholine Receptor $\alpha 7$ (extracellular)	ANC-007