(S)-AMPA

Cat. No.:	HY-100815A				
CAS No.:	83643-88-3				
Molecular Formula:	C ₇ H ₁₀ N ₂ O ₄				
Molecular Weight:	186				
Target:	iGluR				
Pathway:	Membrane Transporter/Ion Channel; Neuronal Signaling				
Storage:	Powder	-20°C	3 years		
	In solvent	-80°C	6 months		
		-20°C	1 month		

SOLVENT & SOLUBILITY

	Mass Solvent Concentration	1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	5.3763 mL	26.8817 mL	53.7634 ml
Stock Solutions	5 mM	1.0753 mL	5.3763 mL	10.7527 ml
	10 mM	0.5376 mL	2.6882 mL	5.3763 mL

DIOLOGICALACTIV	
Description	(S)-AMPA (L-AMPA), an active S-enantiomer of AMPA, is a potent and selective AMPA receptor agonist ^{[1][2]} .
In Vitro	Superfusion of (S)-AMPA (1 μM) significantly attenuates CGRP release in a CB1-dependent manner ^[3] . 24-h exposure to (S)-AMPA (0.01-1000 μM) induces concentration-dependent neuronal cell death (EC ₅₀ of 3 μM) with cellular changes including neurite blebbing, chromatin condensation, and DNA fragmentation, indicative of apoptosis ^[4] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. J A Larm, et al. (S)-5-fluorowillardiine-mediated neurotoxicity in cultured murine cortical neurones occurs via AMPA and kainate receptors. Eur J Pharmacol. 1996 Oct 24;314(1-2):249-54.

[2]. B Ebert, et al. Molecular pharmacology of the AMPA agonist, (S)-2-amino-3-(3-hydroxy-5-phenyl-4-isoxazolyl)propionic acid [(S)-APPA] and the AMPA antagonist, (R)-APPA. Neurochem Int. 1994 Jun;24(6):507-15.

[3]. J W Brooks, et al. (S)-AMPA inhibits electrically evoked calcitonin gene-related peptide (CGRP) release from the rat dorsal horn: reversal by cannabinoid receptor

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 NH_2



antagonist SR141716A. Neurosci Lett. 2004 Nov 30;372(1-2):85-8.

[4]. J A Larm, et al. Apoptosis induced via AMPA-selective glutamate receptors in cultured murine cortical neurons. J Neurochem. 1997 Aug;69(2):617-22.

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

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