CX516

Cat. No.:	HY-10933		
CAS No.:	154235-83-3	3	
Molecular Formula:	$C_{14}H_{15}N_{3}O$		
Molecular Weight:	241.29		
Target:	iGluR		
Pathway:	Membrane Transporter/Ion Channel; Neuronal Signaling		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year

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SOLVENT & SOLUBILITY

In Vitro	H ₂ O : 100 mg/mL (414.44 mM; Need ultrasonic) DMSO : ≥ 41 mg/mL (169.92 mM) * "≥" means soluble, but saturation unknown.						
	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg		
		1 mM	4.1444 mL	20.7220 mL	41.4439 mL		
		5 mM	0.8289 mL	4.1444 mL	8.2888 mL		
		10 mM	0.4144 mL	2.0722 mL	4.1444 mL		
	Please refer to the solubility information to select the appropriate solvent.						
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.75 mg/mL (11.40 mM); Clear solution						
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.75 mg/mL (11.40 mM); Clear solution						
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.75 mg/mL (11.40 mM); Clear solution						

BIOLOGICAL ACTIV	BIOLOGICAL ACTIVITY	
Description	CX516 (BDP 12) is an ampakine and acts as an AMPA receptor positive allosteric modulator for the research of Alzheimer's disease, schizophrenia and mild cognitive impairment (MCI) ^[1] .	
IC ₅₀ & Target	AMPA receptor ^[1]	

Product Data Sheet

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In Vivo	The specific extradimensional deficits produced by sub-chronic or early postnatal postnatal phencyclidine treatment were significantly attenuated by CX516 (10, and 20 mg/kg, s.c.) ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.		
	Animal Model:	Male Lister Hooded rats (56-63 postnatal day) $^{[1]}$	
	Dosage:	5, 10, 20, and 40 mg/kg	
	Administration:	Subcutaneously	
	Result:	Two doses (10 and 20 mg/kg) were highly significant in improving the extradimensional shift deficit induced by either treatment regimes. Two doses (5 and 40 mg/kg) was ineffective at reversing the extradimensional impairment induced by the sub-chronic postnatal phencyclidine treatment regime.	

CUSTOMER VALIDATION

• Toxicol Appl Pharmacol. 2022 Feb 15;439:115924.

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REFERENCES

[1]. Broberg BV, Glenth? j BY, Dias R, et al. Reversal of cognitive deficits by an ampakine (CX516) and sertindole in two animal models of schizophrenia--sub-chronic and early postnatal PCP treatment in attentional set-shifting. Psychopharmacology (Berl). 2009 Nov;206(4):631-40.

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