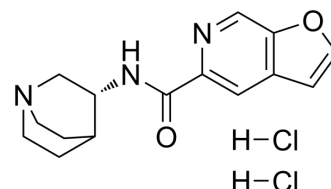


PHA-543613 dihydrochloride

Cat. No.:	HY-105670B
CAS No.:	478148-58-2
Molecular Formula:	C ₁₅ H ₁₉ Cl ₂ N ₃ O ₂
Molecular Weight:	344.24
Target:	nAChR
Pathway:	Membrane Transporter/Ion Channel; Neuronal Signaling
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	PHA-543613 dihydrochloride is a potent, orally active, brain-penetrant and selective α7 nAChR agonist with a K _i value of 8.8 nM. PHA-543613 dihydrochloride displays selectivity for α7-nAChR over α3β4, α1β1γδ, α4β2 and 5-HT3 receptors ^[1] . PHA-543613 dihydrochloride can be used for the cognitive deficits of Alzheimer's disease and schizophrenia research ^{[2][3]} .		
IC ₅₀ & Target	Caution: Product has not been fully validated for medical applications. For research use only. Ki: 8.8 nM (α7 nAChR) ^[1] Tel: 609-228-6898 Fax: 609-228-5909 E-mail: tech@MedChemExpress.com Address: 1 Deer Park Dr, Suite O, Monmouth Junction, NJ 08852, USA		
In Vivo	PHA-543613 dihydrochloride (0.3 mg/kg) successfully reverses Scopolamine-induced short-term memory deficits in rats ^[2] . PHA-543613 dihydrochloride (4 and 12 mg/kg; i.p. once) reduces behavioral deficits and brain edema is dependent on the PI3K-Akt signaling pathway ^[3] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.		
	Animal Model:	Male CD-1 mice with (intracerebral hemorrhage) ICH-induction or sham surgery ^[3]	
	Dosage:	4 and 12 mg/kg	
	Administration:	Intraperitoneal injection; 4 and 12 mg/kg; 1 hour after surgery	
	Result:	Increased p-Akt and decreased p-GSK-3 and CC3 expressions in the ipsilateral hemisphere and reduced the neuronal cell death in the perihematoma area. Attenuated behavioral deficits and brain edema at 72 hours after ICH.	

REFERENCES

- [1]. Donn G Wishka, et al. Discovery of N-[(3R)-1-azabicyclo[2.2.2]oct-3-yl]furo[2,3-c]pyridine-5-carboxamide, an agonist of the α7 nicotinic acetylcholine receptor, for the potential treatment of cognitive deficits in schizophrenia: synthesis and structure-activity relationship. J Med Chem. 2006 Jul 13;49(14):4425-36.
- [2]. Nóra Bruszt, et al. Potentiation of cognitive enhancer effects of Alzheimer's disease medication memantine by α7 nicotinic acetylcholine receptor agonist PHA-543613 in the Morris water maze task. Psychopharmacology (Berl). 2021 Nov;238(11):3273-3281.
- [3]. Krafft PR, et al. α7 nicotinic acetylcholine receptor agonism confers neuroprotection through GSK-3β inhibition in a mouse model of intracerebral hemorrhage. Stroke. 2012 Mar;43(3):844-50.