



上海源叶生物科技有限公司
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产品名称: **MAO-IN-M30 dihydrochloride**

CAS 号: **64821-19-8**

产品货号: **S89887**

生物活性:

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Description	MAO-IN-M30 dihydrochloride is an orally active, brain-permeable, and brain selective irreversible MAO-A (IC50=37 nM) and MAO-B (IC50=57 nM) inhibitor. MAO-IN-M30 dihydrochloride is a potent iron chelator and radical scavenger. MAO-IN-M30 dihydrochloride has a neuroprotective effect against Dexamethasone-induced brain cell apoptosis. MAO-IN-M30 dihydrochloride also exhibits neurorestorative activity in post MPTP and lactacystin models of Parkinson's disease[1][2][3]. MAO-IN-M30 (dihydrochloride) is a click chemistry reagent, it contains an Alkyne group and can undergo copper-catalyzed azide-alkyne cycloaddition (CuAAC) with molecules containing Azide groups.	
IC50 & Target	MAO-A 37 nM (IC50)	MAO-B 57 nM (IC50)
In Vitro	MAO-IN-M30 (0.25 nM; 72 hours) significantly increased cell viability to ~90% after exposure to Dexamethasone[3].	
	MAO-IN-M30 (0-10 μM; 24 hours) enhances PC12 cell survival[4].	
	MAO-IN-M30 treatment significantly decreases the occurrence of fragmented DNA compared to the dexamethasone-treated group in SH-SY5Y cells[3].	
	Cell Viability Assay[3]	
	Cell Line:	SH-SY5Y cells
	Concentration:	0.25 nM
	Incubation Time:	72 hours
	Result:	Significantly increased cell viability to ~ 90% after exposure to Dexamethasone.
	Cell Viability Assay[4]	
	Cell Line:	PC12 cells
Concentration:	0-10 μM	
Incubation Time:	24 hours	
Result:	Enhanced the PC12 cell viability, the cell viability increasing to 85 ± 6 and 90 ± 7%.	
In Vivo	MAO-IN-M30 (0.5-2.5 mg/kg; p.o.; once daily for 14 consecutive days) possesses neuroprotective activities[6].	
	Animal Model:	Male C57/BL mice (20-22 g; MPTP-induced neurotoxicity in mice)[6]
	Dosage:	0.5, 2.5 mg/kg
	Administration:	P.o.; once daily for 14 consecutive days
	Result:	Significantly elevate striatal dopamine levels, reduce its metabolism, and elevate tyrosine-hydroxylase protein levels and activity. Elevated MPTP-reduced dopaminergic and transferrin receptor cell count in the SNpc.
Solvent&Solubility	细胞实验: DMSO 中的溶解度 : 2 mg/mL (6.68 mM; 超声助溶 (<60°C); 吸湿的 DMSO 对产品的溶解度有显著影响, 请使用新开封的 DMSO)	



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	Preparing Stock Solutions	Solvent / Mass Concentration	1 mg	5 mg	10 mg
		1 mM	3.3422 mL	16.7112 mL	33.4225 mL
		5 mM	0.6684 mL	3.3422 mL	6.6845 mL
		* 请根据产品在不同溶剂中的溶解度选择合适的溶剂配制储备液；一旦配成溶液，请分装保存，避免反复冻融造成的产品失效。 储备液的保存方式和期限：-80℃, 6 months; -20℃, 1 month (sealed storage, away from moisture)。-80℃ 储存时，请在 6 个月内使用，-20℃ 储存时，请在 1 个月内使用。			
参考文献	[1]. Gal S, et al. M30, a novel multifunctional neuroprotective drug with potent iron chelating and brain selective monoamine oxidase-ab inhibitory activity for Parkinson's disease. J Neural Transm Suppl. 2006;(70):447-456. [2]. Zheng H, et al. Novel multifunctional neuroprotective iron chelator-monoamine oxidase inhibitor drugs for neurodegenerative diseases: in vitro studies on antioxidant activity, prevention of lipid peroxide formation and monoamine oxidase inhibition. J Neurochem. 2005;95(1):68-78. [3]. Gal S, et al. Novel multifunctional neuroprotective iron chelator-monoamine oxidase inhibitor drugs for neurodegenerative diseases. In vivo selective brain monoamine oxidase inhibition and prevention of MPTP-induced striatal dopamine depletion. J Neurochem. 2005;95(1):79-88. [4]. Gal S, et al. Restoration of nigrostriatal dopamine neurons in post-MPTP treatment by the novel multifunctional brain-permeable iron chelator-monoamine oxidase inhibitor drug, M30. Neurotox Res. 2010;17(1):15-27.				
完整储备液配制表：					
* 请根据产品在不同溶剂中的溶解度选择合适的溶剂配制储备液；一旦配成溶液，请分装保存，避免反复冻融造成的产品失效。 储备液的保存方式和期限：-80℃, 6 months; -20℃, 1 month (sealed storage, away from moisture)。-80℃ 储存时，请在 6 个月内使用，-20℃ 储存时，请在 1 个月内使用。					
可选溶剂	质量 溶剂体积 浓度	1 mg	5 mg	10 mg	25 mg
DMSO	1 mM	3.3422 mL	16.7112 mL	33.4225 mL	83.5561 mL
	5 mM	0.6684 mL	3.3422 mL	6.6845 mL	16.7112 mL