ML 10302 hydrochloride

MedChemExpress

Cat. No.:	HY-14442	
CAS No.:	186826-17-5	^
Molecular Formula:	$C_{15}H_{22}Cl_2N_2O_3$	
Molecular Weight:	349.25	
Target:	5-HT Receptor	H ₂ N O
Pathway:	GPCR/G Protein; Neuronal Signaling	H-CI
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.	

BIOLOGICAL ACTIV					
Description	ML 10302 hydrochloride is a potent and selective 5-HT ₄ receptor agonist, with an EC ₅₀ of 4 nM. ML 10302 hydrochloride displays more than 680-fold selectivity over 5-HT ₃ receptor in binding assay ^{[1][2]} .				
IC ₅₀ & Target	5-HT ₄ Receptor	5-HT ₄ Receptor	5-HT ₃ Receptor		
	4 nM (EC50)	1.07 nM (Ki)	782 nM (Ki)		
In Vitro	ML 10302 hydrochloride shows K _i s of 1.07 nM and 782 nM for 5-HT ₄ receptor and 5-HT ₃ receptor, respectively, in binding assay ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.				
In Vivo	ML 10302 hydrochloride (20 mg/kg; s.c.) induces the soluble form of amyloid precursor protein (sAPP α) production in the cortex of mice ^[3] .				
	Animal Model:	8-weeks old adult male C57BL/6j wild-type mice (23-27 g) ^[3]			
	Dosage:	5 mg/kg, 10 mg/kg, 20 mg/kg			
	Administration:	Subcutaneous injection			
	Result:	Significantly increased sAPPα level in the cortex.			

REFERENCES

[1]. Langlois, M., et al. Design of a potent 5-HT4 receptor agonist with nanomolar affinity. Bioorganic & Medicinal Chemistry Letters. 1994. 4(12), 1433–1436.

[2]. D Yang, et al. New esters of 4-amino-5-chloro-2-methoxybenzoic acid as potent agonists and antagonists for 5-HT4 receptors. J Med Chem. 1997 Feb 14;40(4):608-21.

[3]. M Cachard-Chastel, et al. 5-HT4 receptor agonists increase sAPPα levels in the cortex and hippocampus of male C57BL/6j mice. Br J Pharmacol. 2007 Apr; 150(7): 883–892.

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

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