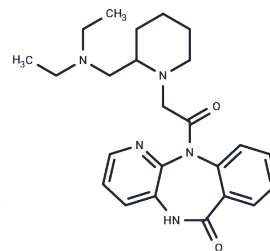


Otenzepad

Chemical Properties

CAS No. :	102394-31-0
Formula:	C ₂₄ H ₃₁ N ₅ O ₂
Molecular Weight:	421.54
Storage:	Powder: -20°C for 3 years In solvent: -80°C for 1 year Actual storage temperature shall be subject to the COA.



Biological Description

Description	Otenzepad (AF-DX 116) is a selective M2 mAChR antagonist with IC ₅₀ values of 386 nM in rat heart and 640 nM in rabbit peripheral lung.
Targets(IC ₅₀)	AChR
In vivo	In rats, Otenzepad (2 mg/kg; s.c.) significantly improved retention relative to vehicle controls. Otenzepad (0.5, 1 mg/kg; s.c.) significantly improved win-stay acquisition[1]. In mice, Otenzepad (0.3, 1.0, or 3.0 mg/kg, i.p.) reverses the effects of insulin on memory and potentiates the effects of glucose[2].

Solubility Information

Solubility	DMSO: 24 mg/mL (56.93 mM), Sonication and heating to 60°C are recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
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Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	2.3723 mL	11.8613 mL	23.7225 mL
5 mM	0.4745 mL	2.3723 mL	4.7445 mL
10 mM	0.2372 mL	1.1861 mL	2.3723 mL
50 mM	0.0474 mL	0.2372 mL	0.4745 mL

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

Reference

Packard MG, et al. Post-training injection of the acetylcholine M2 receptor antagonist AF-DX 116 improves memory. *Brain Res.* 1990 Jul 30;524(1):72-6.

Kopf SR, et al. AF-DX 116, a presynaptic muscarinic receptor antagonist, potentiates the effects of glucose and reverses the effects of insulin on memory. *Neurobiol Learn Mem.* 1998 Nov;70(3):305-13.

Bloom JW, et al. Heterogeneity of the M1 muscarinic receptor subtype between peripheral lung and cerebral cortex demonstrated by the selective antagonist AF-DX 116. *Life Sci.* 1987 Jul 27;41(4):491-6.

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Tel:781-999-4286 E_mail:info@targetmol.com Address:34 Washington Street,Wellesley Hills,MA 02481