



上海源叶生物科技有限公司  
Shanghai yuanye Bio-Technology Co., Ltd  
电话: 021-61312973 传真: 021-55068248  
网址: www.shyuanye.com  
邮箱: shyysw@sina.com

产品名称: 盐酸他利克索

产品别名: **Talipexole dihydrochloride; B-HT 920 dihydrochloride**

生物活性:																											
Description	Talipexole dihydrochloride (B-HT 920 dihydrochloride) is a dopamine D2 receptor agonist, $\alpha$ 2-adrenoceptor agonist and 5-HT3 receptor antagonist, which displays antiParkinsonian activity.																										
IC <sub>50</sub> & Target	IC <sub>50</sub> : 25 nM (Adrenergic receptor $\alpha$ -2, rat)[1].																										
In Vivo	Intravenous injection of 30 micrograms/kg of Talipexole dihydrochloride (B-HT 920) into cats lead initially to an increase in blood pressure and then to a long-lasting decrease in blood pressure and heart rate. Vagally mediated reflex bradycardia elicited by angiotensin injection in beta-adrenoceptor-blocked dogs was facilitated by intracisternal injection of 10 micrograms/kg Talipexole dihydrochloride (B-HT 920).																										
Solvent&Solubility	<p><b>In Vitro:</b> DMSO : 28 mg/mL (99.21 mM; Need ultrasonic and warming)</p> <table border="1"><thead><tr><th rowspan="2">Preparin</th><th rowspan="2">Stock Solutions</th><th>Solvent \ Mass</th><th>1 mg</th><th>5 mg</th><th>10 mg</th></tr><tr><th>Concentration</th><th></th><th></th><th></th></tr></thead><tbody><tr><td>1 mM</td><td>3.5432 mL</td><td>17.7160 mL</td><td>35.4321 mL</td></tr><tr><td>5 mM</td><td>0.7086 mL</td><td>3.5432 mL</td><td>7.0864 mL</td></tr><tr><td>10 mM</td><td>0.3543 mL</td><td>1.7716 mL</td><td>3.5432 mL</td></tr></tbody></table> <p>*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°C, 6 months; -20°C, 1 month。-80°C 储存时,请在 6 个月内使用, -20°C 储存时,请在 1 个月内使用。</p>					Preparin	Stock Solutions	Solvent \ Mass	1 mg	5 mg	10 mg	Concentration				1 mM	3.5432 mL	17.7160 mL	35.4321 mL	5 mM	0.7086 mL	3.5432 mL	7.0864 mL	10 mM	0.3543 mL	1.7716 mL	3.5432 mL
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References	<p>[1]. Ricci and Taira Adrenoceptor involvement in the cardiovascular responses to B-HT 920 in sinoaortic denervated rats. Gen.Pharmacol. (1999)32 29.</p> <p>[2]. Eur J Pharmacol. 1997 May 1;325(2-3):137-44.</p> <p>[3]. Kohno Y, Fukuzaki K, Kitahara K, Koja T.Anti-tremor activity of talipexole produced by selective dopamine D2 receptor stimulation in cynomolgus monkeys with unilateral lesions in the ventromedial tegmentum.Eur J Pharmacol. 1997 Jan 29;319(2-3):197-205.</p> <p>[4]. Momiyama T, Sasa M, Takaori S.Inhibition by talipexole, a thiazolo-azepine derivative, of dopaminergic neurons in the ventral tegmental area.Life Sci. 1991;49(7):535-43.</p> <p>[5]. Nishikawa T, Yamada S, Tsuda A, Tanaka M, Koga I, Uchida Y.Chronic treatment with talipexole dihydrochloride on abnormal involuntary movement in humans.Clin Neuropharmacol. 1990 Jun;13(3):259-63.</p> <p>[6]. Robertson, G.S., et al., In vivo comparisons of the effects of quinpirole and the putative presynaptic dopaminergic agonists B-HT 920 and SND 919 on striatal dopamine and acetylcholine release. J Pharmacol Exp Ther, 1993. 264(3): p. 1344-51.</p>																										