

产品名称：甲磺酸双氢麦角碱

产品别名：**Dihydroergotoxine mesylate; Ergoloid mesylates**

生物活性:																										
Description	Dihydroergotoxine mesylate is a complex of closely related alkaloid salts; Binds with high affinity to the GABAA receptor Cl- channel, producing an allosteric interaction with the benzodiazepine site. IC50 value: Target: Dihydroergotoxine mesylate also interacts with central dopaminergic, serotonergic and adrenergic (α_1) receptors. Dihydroergotoxine mesylate displays antiproliferative activity in vitro (IC50 = 18 - 38 μ M in prostate cancer cells) and exhibits cognition-enhancing, anticonvulsant and sedative activity in vivo.																									
Solvent&Solubility	<p>In Vitro:</p> <p>DMSO : \geq 31 mg/mL (13.39 mM)</p> <p>* "\geq" means soluble, but saturation unknown.</p> <table border="1"><thead><tr><th rowspan="2">Preparing Stock Solutions</th><th>Solvent</th><th>Mass</th><th>Concentration</th><th></th></tr><tr><th></th><th>1 mg</th><th>5 mg</th><th>10 mg</th></tr></thead><tbody><tr><td>1 mM</td><td>0.4320 mL</td><td>2.1601 mL</td><td>4.3201 mL</td></tr><tr><td>5 mM</td><td>0.0864 mL</td><td>0.4320 mL</td><td>0.8640 mL</td></tr><tr><td>10 mM</td><td>0.0432 mL</td><td>0.2160 mL</td><td>0.4320 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。</p> <p>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>					Preparing Stock Solutions	Solvent	Mass	Concentration			1 mg	5 mg	10 mg	1 mM	0.4320 mL	2.1601 mL	4.3201 mL	5 mM	0.0864 mL	0.4320 mL	0.8640 mL	10 mM	0.0432 mL	0.2160 mL	0.4320 mL
Preparing Stock Solutions	Solvent	Mass	Concentration																							
		1 mg	5 mg	10 mg																						
1 mM	0.4320 mL	2.1601 mL	4.3201 mL																							
5 mM	0.0864 mL	0.4320 mL	0.8640 mL																							
10 mM	0.0432 mL	0.2160 mL	0.4320 mL																							
References	<p>[1]. Tvrdeic A, et al. Dihydrogenated ergot compounds bind with high affinity to GABAA receptor-associated Cl- ionophore. Eur J Pharmacol. 1991 Sep 4;202(1):109-11.</p> <p>[2]. Tvrdeic A, et al. Dihydroergotoxine modulation of the GABAA receptor-associated Cl- ionophore in mouse brain. Eur J Pharmacol. 1992 Oct 6;221(1):139-43.</p> <p>[3]. Tvrdeic A, et al. Effect of ergot alkaloids on 3H-flunitrazepam binding to mouse brain GABAA receptors. Coll Antropol. 2003;27 Suppl 1:175-82.</p> <p>[4]. Abdul M, et al. Expression of gamma-aminobutyric acid receptor (subtype A) in prostate cancer. Acta Oncol. 2008;47(8):1546-50.</p>																									