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产品名称: L-655,708

产品别名: L-655708

生物活性:

Description	L-655708 is a potent α 5 subunit-selective GABA _A receptor inverse agonist ($K_i = 0.45$ nM). IC ₅₀ : 0.45 nM (K_i) Target: GABA <i>in vitro</i> : L-655708 is a potent, selective inverse agonist for the benzodiazepine site of GABA _A receptors containing the α 5 subunit ($K_i = 0.45$ nM). Displays 50-100-fold selectivity over GABA _A receptors containing α 1, α 2, α 3 or α 6 subunits in combination with β 3 and γ 2. Enhances LTP in a mouse hippocampal slice model and increases spatial learning, without displaying proconvulsant activity. <i>in vivo</i> : L-655708 at 0.7 mg/kg, administered intraperitoneally, would result in 60-70% occupancy of α 5 GABA _A receptors with limited binding to α 1, α 2, and α 3 subunit-containing GABA _A receptors and no significant off-target behavioral effects, such as sedation and motor impairment.[1]																					
In Vitro: DMSO : 20 mg/mL (58.59 mM; Need ultrasonic) H ₂ O : < 0.1 mg/mL (insoluble)	<table border="1"><thead><tr><th rowspan="2">Preparing 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>2.9295 mL</td><td>14.6473 mL</td><td>29.2946 mL</td></tr><tr><td>5 mM</td><td>0.5859 mL</td><td>2.9295 mL</td><td>5.8589 mL</td></tr><tr><td>10 mM</td><td>0.2929 mL</td><td>1.4647 mL</td><td>2.9295 mL</td></tr></tbody></table>	Preparing Stock Solutions	Solvent / Mass	1 mg	5 mg	10 mg	Concentration				1 mM	2.9295 mL	14.6473 mL	29.2946 mL	5 mM	0.5859 mL	2.9295 mL	5.8589 mL	10 mM	0.2929 mL	1.4647 mL	2.9295 mL
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Solvent&Solubility 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。 In Vivo: 请根据您的实验动物和给药方式选择适当的溶解方案。以下溶解方案都请先按照 <i>In Vitro</i> 方式配制澄清的储备液, 再依次添加助溶剂: ——为保证实验结果的可靠性, 澄清的储备液可以根据储存条件, 适当保存; 体内实验的工作液, 建议您现用现配, 当天使用; 以下溶剂前显示的百分比是指该溶剂在您配制终溶液中的体积占比; 如在配制过程中出现沉淀、析出现象, 可以通过加热和/或超声的方式助溶 1.请依序添加每种溶剂: 10% DMSO → 40% PEG300 → 5% Tween-80 → 45% saline Solubility: ≥ 2 mg/mL (5.86 mM); Clear solution 此方案可获得 ≥ 2 mg/mL (5.86 mM, 饱和度未知) 的澄清溶液。 以 1 mL 工作液为例, 取 100 μL 20.0 mg/mL 的澄清 DMSO 储备液加到 400 μL PEG300 中, 混合均匀, 向上述体系中加入 50 μL Tween-80, 混合均匀; 然后继续加入 450 μL 生理盐水定容至 1 mL。																						
References	[1]. Saab BJ, et al. Short-term memory impairment after isoflurane in mice is prevented by the α 5 γ -aminobutyric acid type A receptor inverse agonist L-655708. Anesthesiology. 2010 Nov;113(5):1061-1071.																					