

产品名称：酒石酸艾芬地尔
产品别名：Ifenprodil tartrate

生物活性：																										
Description	Ifenprodil tartrate is the atypical N-methyl-D-aspartate (NMDA) receptor antagonist, inhibits NMDA-induced currents at NR1A/NR2B receptors with high affinity (IC ₅₀ = 0.34 μM)[1]. Ifenprodil tartrate inhibits G protein-activated inwardly rectifying K ⁺ channels (GIRK, also known as Kir3), reduces inward currents through the basal GIRK activity[2]. Ifenprodil, which is a clinically used cerebral vasodilator, interacts with several receptors, such as alpha1 adrenergic, N-methyl-D-aspartate, serotonin and sigma receptors[2].																									
	IC ₅₀ & Target																									
Solvent&Solubility	<p>In Vitro:</p> <p>DMSO : ≥ 500 mg/mL (1248.47 mM)</p> <p>H₂O : 5 mg/mL (12.48 mM; Need ultrasonic)</p> <p>* "≥" means soluble, but saturation unknown.</p>																									
	<table> <tr> <th rowspan="2"></th><th rowspan="2">Solvent Concentration</th><th colspan="3">Mass</th></tr> <tr> <th>1 mg</th><th>5 mg</th><th>10 mg</th></tr> <tr> <td>Preparing</td><td>1 mM</td><td>2.4969 mL</td><td>12.4847 mL</td><td>24.9694 mL</td></tr> <tr> <td>Stock Solutions</td><td>5 mM</td><td>0.4994 mL</td><td>2.4969 mL</td><td>4.9939 mL</td></tr> <tr> <td></td><td>10 mM</td><td>0.2497 mL</td><td>1.2485 mL</td><td>2.4969 mL</td></tr> </table>		Solvent Concentration	Mass			1 mg	5 mg	10 mg	Preparing	1 mM	2.4969 mL	12.4847 mL	24.9694 mL	Stock Solutions	5 mM	0.4994 mL	2.4969 mL	4.9939 mL		10 mM	0.2497 mL	1.2485 mL	2.4969 mL	<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℃，6 months；-20℃，1 month。-80℃ 储存时，请在 6 个月内使用，-20℃ 储存时，请在 1 个月内使用。</p> <p>In Vivo:</p> <p>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 In Vitro 方式配制澄清的储备液，再依次添加助溶剂：</p> <p>——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</p> <p>1.请依序添加每种溶剂： 1% DMSO→99% saline</p> <p>Solubility: ≥ 1 mg/mL (2.50 mM); Clear solution</p>	
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References	<p>[1]. Williams K, et al. Ifenprodil discriminates subtypes of the N-methyl-D-aspartate receptor: selectivity and mechanisms at recombinant heteromeric receptors. <u>Mol Pharmacol.</u> 1993 Oct;44(4):851-9.</p> <p>[2]. Kobayashi T, et al. Inhibition of G protein-activated inwardly rectifying K⁺ channels by ifenprodil. <u>Neuropsychopharmacology.</u> 2006 Mar;31(3):516-24.</p>																									