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产品名称: **Sunifiram**  
 产品别名: **桑尼非拉姆; DM-235**

生物活性:																	
<b>Description</b>	<p>Sunifiram (DM-235) is a piperazine derived ampakine-like drug which has nootropic effects in animal studies with significantly higher potency than piracetam. IC50 value: Target: in vitro: DM 232 and DM 235 are novel anti-amnesic compounds structurally related to ampakines. The involvement of AMPA receptors in the mechanism of action of DM 232 and DM 235 was, therefore, investigated in vivo and in vitro. Both compounds (0.1 mg/kg i.p.) were able to reverse the amnesia induced by the AMPA receptor antagonist NBQX (30 mg/kg i.p.) in the mouse passive avoidance test. At the effective doses, the investigated compounds did not impair motor coordination, as revealed by the rota rod test, nor modify spontaneous motility and inspection activity, as revealed by the hole board test [1]. In mouse hippocampal slices, sunifiram at 10-100 nM significantly enhanced LTP in a bell-shaped dose-response relationship which peaked at 10 nM. The enhancement of LTP by sunifiram treatment was inhibited by 7-chloro-kynurenic acid (7-ClKN), an antagonist for glycine-binding site of NMDAR, but not by ifenprodil, an inhibitor for polyamine site of NMDAR [2]. in vivo: OBX mice were administered once a day for 7-12 days with sunifiram (0.01-1.0 mg/kg p.o.) from 10 days after operation with or without gavestinel (10 mg/kg i.p.), which is glycine-binding site inhibitor of N-methyl-D-aspartate receptor (NMDAR) [3].</p>																
<b>Solvent&amp;Solubility</b>	<p><b>In Vitro:</b>            H<sub>2</sub>O : 50 mg/mL (203.00 mM; Need ultrasonic)            DMSO : ≥ 2.6 mg/mL (10.56 mM)            * "≥" means soluble, but saturation unknown.</p>																
	<b>Preparing</b>	<table border="1"> <thead> <tr> <th style="text-align: center;">Solvent</th> <th style="text-align: center;">Mass</th> <th style="text-align: center;">Concentration</th> <th style="text-align: center;">1 mg</th> <th style="text-align: center;">5 mg</th> <th style="text-align: center;">10 mg</th> </tr> </thead> <tbody> <tr> <td></td> <td></td> <td>1 mM</td> <td>4.0601 mL</td> <td>20.3004 mL</td> <td>40.6009 mL</td> </tr> </tbody> </table>	Solvent	Mass	Concentration	1 mg	5 mg	10 mg			1 mM	4.0601 mL	20.3004 mL	40.6009 mL			
	Solvent	Mass	Concentration	1 mg	5 mg	10 mg											
			1 mM	4.0601 mL	20.3004 mL	40.6009 mL											
<b>Stock Solutions</b>		5 mM	0.8120 mL	4.0601 mL	8.1202 mL												
		10 mM	0.4060 mL	2.0300 mL	4.0601 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。            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>																	
<b>References</b>	<p>[1]. Galeotti N, et al. AMPA-receptor activation is involved in the anti-amnesic effect of DM 232 (unifiram) and DM 235 (sunifiram). Naunyn Schmiedebergs Arch Pharmacol. 2003 Dec;368(6):538-45.            [2]. Moriguchi S, et al. Novel nootropic drug sunifiram enhances hippocampal synaptic efficacy via glycine-binding site of N-methyl-D-aspartate receptor. Hippocampus. 2013 Jun 3.            [3]. Moriguchi S, et al. Novel nootropic drug sunifiram improves cognitive deficits via CaM kinase II and protein kinase C activation in olfactory bulbectomized mice. Behav Brain Res. 2013 Apr 1;242:150-7.</p>																