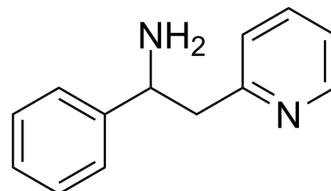


## (Rac)-Lanicemine

<b>Cat. No.:</b>	HY-108235B		
<b>CAS No.:</b>	61890-25-3		
<b>Molecular Formula:</b>	C <sub>13</sub> H <sub>14</sub> N <sub>2</sub>		
<b>Molecular Weight:</b>	198.26		
<b>Target:</b>	iGluR		
<b>Pathway:</b>	Membrane Transporter/Ion Channel; Neuronal Signaling		
<b>Storage:</b>	Pure form	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



### SOLVENT & SOLUBILITY

<b>In Vitro</b>	DMSO : 100 mg/mL (504.39 mM; Need ultrasonic)			
		Solvent Concentration	Mass	
			1 mg	5 mg
			10 mg	
<b>Preparing Stock Solutions</b>	<b>1 mM</b>	5.0439 mL	25.2194 mL	50.4388 mL
	<b>5 mM</b>	1.0088 mL	5.0439 mL	10.0878 mL
	<b>10 mM</b>	0.5044 mL	2.5219 mL	5.0439 mL
Please refer to the solubility information to select the appropriate solvent.				
<b>In Vivo</b>	<ol style="list-style-type: none"> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 40% PEG300 &gt;&gt; 5% Tween-80 &gt;&gt; 45% saline Solubility: ≥ 2.5 mg/mL (12.61 mM); Clear solution</li> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (12.61 mM); Clear solution</li> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 90% corn oil Solubility: ≥ 2.5 mg/mL (12.61 mM); Clear solution</li> </ol>			

### BIOLOGICAL ACTIVITY

<b>Description</b>	(Rac)-Lanicemine ((Rac)-AZD6765) is the racemate of Lanicemine. Lanicemine (AZD6765) is a low-trapping NMDA channel blocker (K <sub>i</sub> of 0.56-2.1 μM for NMDA receptor; IC <sub>50</sub> s of 4-7 μM and 6.4 μM in CHO and Xenopus oocyte cells, respectively). Antidepressant effects <sup>[1]</sup> .
<b>IC<sub>50</sub> &amp; Target</b>	NMDA receptor <sup>[1]</sup>
<b>In Vivo</b>	Lanicemine produces sustained antidepressant efficacy with minimal psychotomimetic adverse effects <sup>[1]</sup> . Lanicemine (3, 10

or 30 mg/kg; intraperitoneal) not only engages brain circuits involved in the generation of gamma- electroencephalography (EEG), but also influences these networks independent of the broader systems-level disruptions typical of ketamine<sup>[1]</sup>. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Male Sprague-Dawley rats <sup>[1]</sup>
Dosage:	3, 10 or 30 mg/kg
Administration:	Intraperitoneal
Result:	Produced pronounced dose-dependent elevations in spontaneous gamma-band EEG, but only gamma changes for Ketamine were tightly coupled to increases in locomotor activity.

## REFERENCES

[1]. Sanacora G, et al. Lanicemine: a low-trapping NMDA channel blocker produces sustained antidepressant efficacy with minimal psychotomimetic adverse effects. *Mol Psychiatry*. 2014 Sep;19(9):978-85.

**Caution: Product has not been fully validated for medical applications. For research use only.**

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