



# SZABO SCANDIC

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### SZABO-SCANDIC HandelsgmbH

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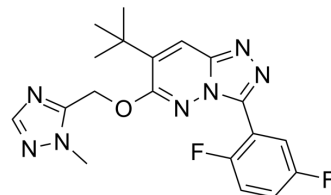
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## L-838417

<b>Cat. No.:</b>	HY-W009009		
<b>CAS No.:</b>	286456-42-6		
<b>Molecular Formula:</b>	C <sub>19</sub> H <sub>19</sub> F <sub>2</sub> N <sub>7</sub> O		
<b>Molecular Weight:</b>	399.41		
<b>Target:</b>	GABA Receptor		
<b>Pathway:</b>	Membrane Transporter/Ion Channel; Neuronal Signaling		
<b>Storage:</b>	Powder	-20°C	3 years
	In solvent	-80°C	6 months
		-20°C	1 month



### SOLVENT & SOLUBILITY

<b>In Vitro</b>	DMSO : 25 mg/mL (62.59 mM; Need ultrasonic)					
	<b>Preparing Stock Solutions</b>	<b>Solvent</b>	<b>Mass</b>	<b>1 mg</b>	<b>5 mg</b>	<b>10 mg</b>
		<b>Concentration</b>				
		<b>1 mM</b>		2.5037 mL	12.5185 mL	25.0369 mL
		<b>5 mM</b>		0.5007 mL	2.5037 mL	5.0074 mL
	<b>10 mM</b>		0.2504 mL	1.2518 mL	2.5037 mL	
Please refer to the solubility information to select the appropriate solvent.						
<b>In Vivo</b>	1. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2 mg/mL (5.01 mM); Clear solution					

### BIOLOGICAL ACTIVITY

<b>Description</b>	L-838417 is a selective partial agonist at the α <sub>2</sub> , α <sub>3</sub> and α <sub>5</sub> subtypes of the GABA <sub>A</sub> receptor and an antagonist at the α <sub>1</sub> , with binding K <sub>i</sub> values of 0.79 nM, 0.67 nM, 1.67 nM, 267 nM, 2.25 nM and 2183 nM for α <sub>1</sub> β <sub>3</sub> γ <sub>2</sub> , α <sub>2</sub> β <sub>3</sub> γ <sub>2</sub> , α <sub>3</sub> β <sub>3</sub> γ <sub>2</sub> , α <sub>4</sub> β <sub>3</sub> γ <sub>2</sub> , α <sub>5</sub> β <sub>3</sub> γ <sub>2</sub> and α <sub>6</sub> β <sub>3</sub> γ <sub>2</sub> <sup>[1]</sup> .
<b>IC<sub>50</sub> &amp; Target</b>	Ki: 0.79 nM (α <sub>1</sub> β <sub>3</sub> γ <sub>2</sub> ), 0.67 nM (α <sub>2</sub> β <sub>3</sub> γ <sub>2</sub> ), 1.67 nM (α <sub>3</sub> β <sub>3</sub> γ <sub>2</sub> ), 267 nM (α <sub>4</sub> β <sub>3</sub> γ <sub>2</sub> ), 2.25 nM (α <sub>5</sub> β <sub>3</sub> γ <sub>2</sub> ) and 2183 nM (α <sub>6</sub> β <sub>3</sub> γ <sub>2</sub> ) <sup>[1]</sup> .
<b>In Vivo</b>	L-838417 (1.0 mg/kg) produces anxiolytic effects in adult rat, as indexed by a transformation of social avoidance into preference and an increase in social investigation <sup>[2]</sup> . L-838417 (2.0 mg/kg) eliminates social avoidance, but has no anxiolytic effects on social investigation <sup>[2]</sup> . L-838417 (0.5 mg/kg) reverses the anxiogenic effects of prior stress regardless of age, but with doses ≥ 1 mg/kg decreases social investigation, an effect possibly due in part to locomotor-impairing effects of this compound <sup>[2]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

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Animal Model:	Male and female adolescent and adult Sprague–Dawley rats <sup>[2]</sup> .
Dosage:	0, 0.5, 1.0, 2.0, or 4.0 mg/kg.
Administration:	IP.
Result:	Adolescents required a higher dose (2 mg/kg) to attenuate their social avoidance. The lowest dose of 0.5 mg/kg was sufficient to reverse the anxiogenic effects of repeated restraint as reflected by a significant increase in the coefficient relative to vehicle-treated animals.

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## REFERENCES

- [1]. Ciara McCabe, et al. Subtype-selective GABAergic drugs facilitate extinction of mouse operant behavior. *Neuropharmacology*. 2004 Feb;46(2):171-8.
- [2]. Melissa Morales, et al. Anxiolytic effects of the GABAA receptor partial agonist, L-838,417: Impact of age, test context familiarity, and stress. *Pharmacol Biochem Behav*. 2013 Aug;109:31-7.
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**Caution: Product has not been fully validated for medical applications. For research use only.**

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