# Inhibitors

# **SNC162**

Cat. No.: HY-107741 CAS No.: 178803-51-5 Molecular Formula:  $C_{27}H_{37}N_{3}O$ Molecular Weight: 419.6

Target: **Opioid Receptor** 

Pathway: GPCR/G Protein; Neuronal Signaling

Storage: Powder -20°C 3 years

> In solvent -80°C 6 months

> > -20°C 1 month

**Product** Data Sheet

## **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 5 mg/mL (11.92 mM; ultrasonic and warming and heat to 60°C)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.3832 mL	11.9161 mL	23.8322 mL
	5 mM	0.4766 mL	2.3832 mL	4.7664 mL
	10 mM	0.2383 mL	1.1916 mL	2.3832 mL

Please refer to the solubility information to select the appropriate solvent.

## **BIOLOGICAL ACTIVITY**

Description	SNC162 is a delta-opioid receptor agonist with an IC <sub>50</sub> of 0.94 nM. SNC162 has antidepressant-like effects and produces a selective enhancement of the antinociceptive effects of fentanyl in rhesus monkeys <sup>[1][2]</sup> .
IC <sub>50</sub> & Target	${ m IC}_{50}$ : 0.94 nM (delta-opioid receptor) $^{[1]}$

### **REFERENCES**

[1]. Jutkiewicz EM, et al. Delta-opioid agonists: differential efficacy and potency of SNC80, its 3-OH (SNC86) and 3-desoxy (SNC162) derivatives in Sprague-Dawley rats. J Pharmacol Exp Ther. 2004 Apr;309(1):173-81.

[2]. Banks ML, et al. Selective enhancement of fentanyl-induced antinociception by the delta agonist SNC162 but not by ketamine in rhesus monkeys: Further evidence supportive of delta agonists as candidate adjuncts to mu opioid analgesics. Pharmacol Biochem Be

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$ 

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