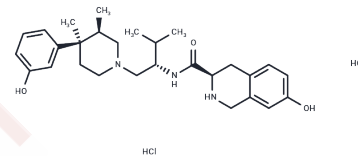


## JDTic Dihydrochloride

## Chemical Properties

CAS No. :	785835-79-2
Formula:	C <sub>28</sub> H <sub>41</sub> Cl <sub>2</sub> N <sub>3</sub> O <sub>3</sub>
Molecular Weight:	538.55
Storage:	Store at low temperature Powder: -20°C for 3 years   In solvent: -80°C for 1 year <i>Actual storage temperature shall be subject to the COA.</i>



## Biological Description

Description	JDTic Dihydrochloride is a high-affinity and selective $\kappa$ -opioid receptor (KOR) antagonist that blocks dynorphin-KOR signalling, exhibiting antidepressant, anxiolytic, and anti-relapse effects against cocaine and nicotine.
Targets(IC50)	Opioid Receptor
In vivo	<p><b>Methods:</b> In a hangover-induced anxiety rat model, JDTic Dihydrochloride was administered via intraperitoneal injection at 3 mg/kg to assess anxiety-like behavior and at 10 mg/kg to evaluate alcohol-related behaviors, with a 2-hour pretreatment period.</p> <p><b>Results:</b> JDTic Dihydrochloride (3 mg/kg, ip) significantly reversed anxiety-like behavior in hangover rats; the 10 mg/kg dose reduced alcohol self-administration, suppressed cue-induced reinstatement of alcohol seeking, and specifically blocked the effects of KOR agonists after 2 hours of pretreatment. [2]</p>

## Solubility Information

Solubility	DMSO: 80 mg/mL (148.55 mM), Sonication is recommended. H <sub>2</sub> O: 40 mg/mL (74.27 mM), Sonication is recommended. ( $< 1$ mg/ml refers to the product slightly soluble or insoluble)
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### Preparing Stock Solutions

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	1mg	5mg	10mg
1 mM	1.8568 mL	9.2842 mL	18.5684 mL
5 mM	0.3714 mL	1.8568 mL	3.7137 mL
10 mM	0.1857 mL	0.9284 mL	1.8568 mL
50 mM	0.0371 mL	0.1857 mL	0.3714 mL

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Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

### Reference

Jackson KJ, et al. Effect of the selective kappa-opioid receptor antagonist JD<sub>Tic</sub> on nicotine antinociception, reward, and withdrawal in the mouse. *Psychopharmacology (Berl)*. 2010 Jun;210(2):285-94.

Schank JR, et al. The kappa opioid receptor antagonist JD<sub>Tic</sub> attenuates alcohol seeking and withdrawal anxiety. *Addict Biol*. 2012 May;17(3):634-47. doi: 10.1111/j.1369-1600.2012.00455.x.

Beardsley PM, et al. Effectiveness of analogs of the kappa opioid receptor antagonist (3R)-7-hydroxy-N-((1S)-1-(((3R,4R)-4-(3-hydroxyphenyl)-3,4-dimethyl-1-piperidinyl)methyl)-2-methylpropyl)-1,2,3,4-tetrahydro-3-isoquinolinecarboxamide (JD<sub>Tic</sub>) to reduce

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