**Proteins** 

# **Product** Data Sheet

## PF-5006739

Cat. No.: HY-12443 CAS No.: 1293395-67-1 Molecular Formula:  $C_{22}H_{22}FN_{7}O$ Molecular Weight: 419.45

Target: Casein Kinase

Pathway: Cell Cycle/DNA Damage; Stem Cell/Wnt

4°C, protect from light Storage:

\* In solvent: -80°C, 6 months; -20°C, 1 month (protect from light)

### **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 100 mg/mL (238.41 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.3841 mL	11.9204 mL	23.8407 mL
	5 mM	0.4768 mL	2.3841 mL	4.7681 mL
	10 mM	0.2384 mL	1.1920 mL	2.3841 mL

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

In Vivo

- 1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (5.96 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (5.96 mM); Clear solution

### **BIOLOGICAL ACTIVITY**

Description PF-5006739 is a potent and selective inhibitor of CK1 $\delta$ / $\epsilon$  with IC $_{50}$ s of 3.9 nM and 17.0 nM, respectively. PF-5006739 is a potential therapeutic agent for a range of psychiatric disorders with low nanomolar in vitro potency for  $CK1\delta/\epsilon$  and high kinome selectivity. PF-5006739 attenuats opioid agent-seeking behavior in a rodent operant reinstatement model in animals in a dose-dependent manner [1]. PF-5006739 improves glucose tolerance in both diet-induced obesity (DIO) and genetic (ob/ob) mice models of obesity<sup>[2]</sup>.

IC<sub>50</sub> & Target

CK1δ 3.9 nM (IC<sub>50</sub>)

#### **REFERENCES**

[1]. Wager TT, et al. Casein kinase 1δ/ε inhibitor PF-5006739 attenuates opioid drug-seeking behavior. ACS Chem Neurosci. 2014 Dec 17;5(12):1253-65. [2]. Cunningham PS, et al. Targeting of the circadian clock via CK1δ/ε to improve glucose homeostasis in obesity. Sci Rep. 2016 Jul 21;6:29983.					
Caution: Product has not been fully validated for medical applications. For research use only.					
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