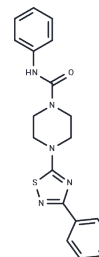


JNJ-1661010

Chemical Properties

CAS No. : 681136-29-8
Formula: C₁₉H₁₉N₅O₂
Molecular Weight: 365.45
Storage: Powder: -20°C for 3 years | In solvent: -80°C for 1 year
Actual storage temperature shall be subject to the COA.



Biological Description

Description	JNJ-1661010 (Takeda-25) is an effective and specific FAAH inhibitor (IC ₅₀ : 10/ 12 nM for rat/human), shows >100-fold selectivity for FAAH-1 than FAAH-2.
Targets(IC ₅₀)	FAAH
In vitro	Preincubation of JNJ-1661010 with fatty acid amide hydrolase showed a slow reversible interaction between JNJ-1661010 and the active site, which was catalyzed by high temperature. In the hydrophobic channel, JNJ-1661010 could dock with phenylthiadiazole, and in the hydrophilic pocket of fatty acid amide hydrolase, it docked with phenylurea.
In vivo	Preincubation of JNJ-1661010 with fatty acid amide hydrolase showed a slow reversible interaction between JNJ-1661010 and the active site, which was catalyzed by high temperature. In the hydrophobic channel, JNJ-1661010 could dock with phenylthiadiazole, and in the hydrophilic pocket of fatty acid amide hydrolase, it docked with phenylurea.

Solubility Information

Solubility	Ethanol: 3.7 mg/mL (10.12 mM), Sonication is recommended. DMSO: 36.6 mg/mL (100.15 mM), Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
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Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	2.7364 mL	13.6818 mL	27.3635 mL
5 mM	0.5473 mL	2.7364 mL	5.4727 mL
10 mM	0.2736 mL	1.3682 mL	2.7364 mL
50 mM	0.0547 mL	0.2736 mL	0.5473 mL

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

- Karbarz MJ, et al. *Anesth Analg*, 2009, 108(1), 316-329.
- Keith JM, et al. *Bioorg Med Chem Lett*, 2008, 18(17), 4838-4843.
- Palumbo-Zerr K, et al. *Ann Rheum Dis*, 2012 Aug 22.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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