

## BMS-470539 dihydrochloride

## Chemical Properties

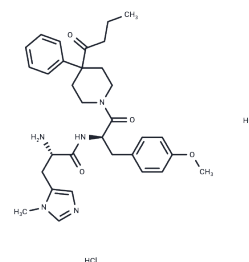
CAS No. : 2341796-82-3

Formula: C<sub>32</sub>H<sub>43</sub>Cl<sub>2</sub>N<sub>5</sub>O<sub>4</sub>

Molecular Weight: 632.62

Storage: Store at low temperature, Keep away from moisture  
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	BMS-470539 dihydrochloride is a selective and highly potent melanocortin 1 receptor (MC-1 R) agonist with anti-inflammatory activity. BMS-470539 attenuates oxidative stress and neuronal apoptosis via the MC1R/cAMP/PKA/Nurr1 signaling pathway in a neonatal hypoxia-ischemia rat model.
Targets(IC50)	Melanocortin Receptor
In vitro	<b>METHODS:</b> BMS-470539 dihydrochloride (0-1 μM, 15 minutes) was used to treat HBL cells stably transfected with NF-κB luciferase reporter gene, and then treated with 0.5 ng/mL TNF-α for 4 hours to study its effect on stable transfection. Effects of NF-κB luciferase reporter gene on TNF-α-stimulated luciferase activation in HBL cells. <b>RESULTS:</b> BMS-470539 dihydrochloride treatment of HBL cells with NF-κB luciferase reporter gene caused a dose-dependent significant reduction in TNF-α-stimulated NF-κB luciferase activity. [2]
In vivo	<b>METHODS:</b> BMS-470539 dihydrochloride (2.05-18.47 mg/kg, intravenous injection, 125 minutes) was used to treat WT and MC1 receptor-negative e/e mice to study its effects on mesenteric microcirculation. <b>RESULTS:</b> BMS-470539 dihydrochloride treatment of mice inhibited cell adhesion and migration, had no effect on cell rolling, and also inhibited the tissue expression of two chemokines, CXCL1 and CCL2. [3]

### Preparing Stock Solutions

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	<b>1mg</b>	<b>5mg</b>	<b>10mg</b>
1 mM	1.5807 mL	7.9036 mL	15.8073 mL
5 mM	0.3161 mL	1.5807 mL	3.1615 mL
10 mM	0.1581 mL	0.7904 mL	1.5807 mL
50 mM	0.0316 mL	0.1581 mL	0.3161 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

- Herpin TF, et al. Discovery of tyrosine-based potent and selective melanocortin-1 receptor small-molecule agonists with anti-inflammatory properties. *J Med Chem.* 2003 Mar 27;46(7):1123-6.
- Kang L, et al. A selective small molecule agonist of the melanocortin-1 receptor inhibits lipopolysaccharide-induced cytokine accumulation and leukocyte infiltration in mice. *J Leukoc Biol.* 2006 Oct;80(4):897-904. Epub 2006 Aug 3.
- Leoni G, et al. The melanocortin MC(1) receptor agonist BMS-470539 inhibits leucocyte trafficking in the inflamed vasculature. *Br J Pharmacol.* 2010 May;160(1):171-80.

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