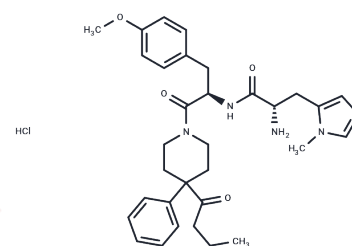


## BMS-470539 dihydrochloride

## Chemical Properties

|                   |                                                          |
|-------------------|----------------------------------------------------------|
| CAS No. :         | 2341796-82-3                                             |
| Formula:          | C32H42ClN5O4                                             |
| Molecular Weight: | 596.17                                                   |
| Appearance:       | no data available                                        |
| Storage:          | Powder: -20°C for 3 years   In solvent: -80°C for 1 year |



## Biological Description

|               |                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   |
|---------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Description   | BMS-470539 dihydrochloride is a highly potent and selective agonist of the melanocortin-1 receptor (MC-1R; IC50: 120 nM; EC50: 28 nM) with anti-inflammatory properties. It does not activate MC-3R and is a very weak partial agonist at MC-4R and MC-5R.                                                                                                                                                                                                                                                                                                                        |
| Targets(IC50) | Others                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            |
| In vitro      | An established HBL melanoma cell line stably expresses an NF-κB luciferase reporter, allowing for the observation that TNF-α at 0.5 ng/mL incites a dose-dependent escalations in NF-κB luciferase activity. Administering BMS-470539 to these HBL-NF-κB cells leads to a dose-dependent diminution in TNF-α-induced NF-κB luciferase activity, while it remains ineffective in altering luciferase reporter activity in the absence of TNF-α induction. In nontransfected HBL cells, BMS-470539 administration shows a dose-dependent deterrence of NF-κB nuclear translocation. |
| In vivo       | The treatment of BMS-470539 (2.05-18.47 mg/kg; i.v.; for 125 minutes; WT and MC1 receptor recessive e/e mice) inhibits cell emigration and adhesion with no effect on cell rolling. It also inhibits the tissue expression of CXCL1/CCL2 [3].                                                                                                                                                                                                                                                                                                                                     |

## Preparing Stock Solutions

|       | 1mg       | 5mg       | 10mg       |
|-------|-----------|-----------|------------|
| 1 mM  | 1.6774 mL | 8.3869 mL | 16.7737 mL |
| 5 mM  | 0.3355 mL | 1.6774 mL | 3.3547 mL  |
| 10 mM | 0.1677 mL | 0.8387 mL | 1.6774 mL  |
| 50 mM | 0.0335 mL | 0.1677 mL | 0.3355 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.

## 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.

**Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins**

This product is for Research Use Only· Not for Human or Veterinary or Therapeutic Use

Tel:781-999-4286    E\_mail:info@targetmol.com    Address:36 Washington Street,Wellesley Hills,MA 02481