

Mirogabalin besylate

Chemical Properties

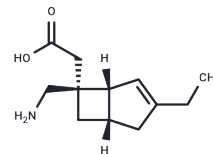
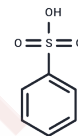
CAS No. : 1138245-21-2

Formula: C₁₈H₂₅NO₅S

Molecular Weight: 367.46

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 | Mirogabalin besylate is a potent and orally active ligand that selectively targets the $\alpha 2\delta$ subunit of voltage-gated calcium channels. It displays high affinity (K_d) values, namely 13.5 nM, 22.7 nM, 27 nM, and 47.6 nM, for human $\alpha 2\delta$ -1, human $\alpha 2\delta$ -2, rat $\alpha 2\delta$ -1, and rat $\alpha 2\delta$ -2, respectively. |
| Targets(IC50) | Others, Calcium Channel |
| In vitro | Mirogabalin besylate specifically targets the $\alpha 2\delta$ subunit of voltage-gated calcium channels, demonstrating varying binding affinities with dissociation constants (K_d) of 13.5 nM for human $\alpha 2\delta$ -1, 22.7 nM for human $\alpha 2\delta$ -2, 27 nM for rat $\alpha 2\delta$ -1, and 47.6 nM for rat $\alpha 2\delta$ -2. Additionally, it has shown a strong binding affinity to the gabapentin site in rat cortical brain homogenates, indicated by an IC ₅₀ value of 16.0 nM. Notably, at a concentration of 50 μ M, mirogabalin does not impact any other receptors, channels, transporters, or enzymes [1]. |
| In vivo | Mirogabalin besylate, administered at doses of 3 and 10 mg/kg, significantly increases area under the curve (AUC) values for 0-8 hours in a dose-dependent manner in rats with partial sciatic nerve ligation. Additionally, doses of 2.5, 5, and 10 mg/kg of Mirogabalin significantly elevate AUC values for 0-12 hours, demonstrating dose-dependent analgesic enhancement, with estimated effective doses (ED) of 4.4 mg/kg on day 1, 3.1 mg/kg on day 3, and less than 2.5 mg/kg by day 5. Furthermore, at 3 and 10 mg/kg, Mirogabalin besylate does not significantly impact rota-rod performance or locomotor activity when administered orally but does show significant inhibition of rota-rod performance at higher doses of 10, 30, and 100 mg/kg, as well as decreases in locomotor activity at 30 and 100 mg/kg in rats [1]. |

Preparing Stock Solutions

| | 1mg | 5mg | 10mg |
|-------|------------|------------|-------------|
| 1 mM | 2.7214 mL | 13.6069 mL | 27.2138 mL |
| 5 mM | 0.5443 mL | 2.7214 mL | 5.4428 mL |
| 10 mM | 0.2721 mL | 1.3607 mL | 2.7214 mL |
| 50 mM | 0.0544 mL | 0.2721 mL | 0.5443 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.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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

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