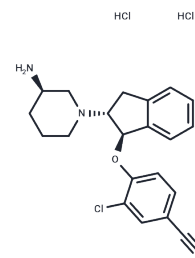


SAR7334 hydrochloride

Chemical Properties

| | |
|-------------------|--|
| CAS No. : | 1333207-63-8 |
| Formula: | C ₂₁ H ₂₄ Cl ₃ N ₃ O |
| Molecular Weight: | 440.79 |
| Storage: | Store at low temperature Powder: -20°C for 3 years In solvent: -80°C for 1 year <i>Actual storage temperature shall be subject to the COA.</i> |



Biological Description

| | |
|----------------------------|---|
| Description | SAR7334 hydrochloride, a potent and specific inhibitor of TRPC6 (IC ₅₀ of 7.9 nM), effectively targets and inhibits the TRPC6 channel. |
| Targets(IC ₅₀) | Others,TRP/TRPV Channel |
| In vitro | SAR7334 inhibits TRPC6, TRPC3, and TRPC7-mediated Ca ²⁺ influx into cells (IC ₅₀ s of 9.5, 282, and 226 nM, respectively) [1][2][3], while TRPC4 and TRPC5-mediated Ca ²⁺ entry remains unaffected. At 1 μM, SAR7334 significantly blocks Ang II-evoked calcium influx in podocytes [1]. SAR7334 dose-dependently reduces TRPC6 currents with an IC ₅₀ of 7.9 nM, and at 100 nM, it substantially reduces TRPC6 currents [3]. |
| In vivo | In isolated perfused lungs from mice, SAR7334 (10?mg/kg, p.o.) inhibits TRPC6-dependent acute HPV. it is suitable for chronic oral administration. In an initial short-term study, SAR7334 does not change mean arterial pressure in spontaneously hypertensive rats (SHR)[3]. |

Solubility Information

| | |
|---------------------|--|
| Solubility | DMSO: 100 mg/mL (226.87 mM),Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble) |
| In vivo Formulation | 10% DMSO+40% PEG300+5% Tween 80+45% Saline: 4 mg/mL (9.07 mM),Sonication is recommended. <i>Please add the solvents sequentially, clarifying the solution as much as possible before adding the next one. Dissolve by heating and/or sonication if necessary. Working solution is recommended to be prepared and used immediately. The formulation provided above is for reference purposes only. In vivo formulations may vary and should be modified based on specific experimental conditions.</i> |

Preparing Stock Solutions

| | 1mg | 5mg | 10mg |
|-------|-----------|------------|------------|
| 1 mM | 2.2687 mL | 11.3433 mL | 22.6865 mL |
| 5 mM | 0.4537 mL | 2.2687 mL | 4.5373 mL |
| 10 mM | 0.2269 mL | 1.1343 mL | 2.2687 mL |
| 50 mM | 0.0454 mL | 0.2269 mL | 0.4537 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

Ilatovskaya DV, et al. The Role of Angiotensin II in Glomerular Volume Dynamics and Podocyte Calcium Handling. *Sci Rep.* 2017 Mar 22;7(1):299.

Chauvet S, et al. Pharmacological Characterization of the Native Store-Operated Calcium Channels of Cortical Neurons from Embryonic Mouse Brain. *Front Pharmacol.* 2016 Dec 12;7:486.

Maier T, et al. Discovery and pharmacological characterization of a novel potent inhibitor of diacylglycerol-sensitive TRPC cation channels. *Br J Pharmacol.* 2015 Jul;172(14):3650-60.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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