

SAR7334

## Chemical Properties

CAS No. : 1333210-07-3

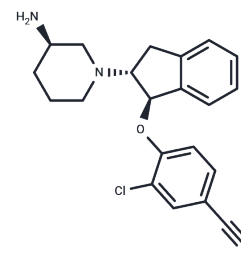
Formula: C<sub>21</sub>H<sub>22</sub>ClN<sub>3</sub>O

Molecular Weight: 367.87

Store at low temperature

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	SAR7334 (TRCP6-IN-1) is a potent and specific inhibitor of TRPC6 (IC <sub>50</sub> of 7.9 nM).
Targets (IC <sub>50</sub> )	TRP/TRPV Channel
In vitro	SAR7334 is an inhibitor of TRPC6, TRPC3 and TRPC7-mediated Ca <sup>2+</sup> influx into cells (IC <sub>50</sub> s of 9.5, 282 and 226 nM)[1][2][3], whereas TRPC4 and TRPC5-mediated Ca <sup>2+</sup> entry is not affected. SAR7334 (1 μM) results in a major block of the Ang II-evoked calcium influx in the podocytes[1]. SAR7334 dose-dependently reduces TRPC6 currents (IC <sub>50</sub> of 7.9 nM). SAR7334 (100 nM) substantially reduces TRPC6 currents[3].
In vivo	In isolated perfused lungs from mice, SAR7334 (10mg/kg, p.o.) inhibits TRPC6-dependent acute HPV. SAR7334 demonstrates that 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: 350 mg/mL (951.42 mM), Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
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### Preparing Stock Solutions

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	1mg	5mg	10mg
1 mM	2.7184 mL	13.5918 mL	27.1835 mL
5 mM	0.5437 mL	2.7184 mL	5.4367 mL
10 mM	0.2718 mL	1.3592 mL	2.7184 mL
50 mM	0.0544 mL	0.2718 mL	0.5437 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

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.

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