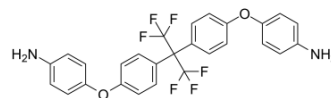


## GI-530159

<b>Cat. No.:</b>	HY-W013712
<b>CAS No.:</b>	69563-88-8
<b>Molecular Formula:</b>	C <sub>27</sub> H <sub>20</sub> F <sub>6</sub> N <sub>2</sub> O <sub>2</sub>
<b>Molecular Weight:</b>	518.45
<b>Target:</b>	Potassium Channel
<b>Pathway:</b>	Membrane Transporter/Ion Channel
<b>Storage:</b>	4°C, sealed storage, away from moisture and light * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture and light)



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : 250 mg/mL (482.21 mM; Need ultrasonic)

Concentration	Mass		
	1 mg	5 mg	10 mg
1 mM	1.9288 mL	9.6441 mL	19.2883 mL
5 mM	0.3858 mL	1.9288 mL	3.8577 mL
10 mM	0.1929 mL	0.9644 mL	1.9288 mL

Please refer to the solubility information to select the appropriate solvent.

### BIOLOGICAL ACTIVITY

#### Description

GI-530159 is a selective, mechanosensitive opener of TREK1 (K<sub>2p</sub>2.1) and TREK2 (K<sub>2p</sub>10.1) channels, with an EC<sub>50</sub> of 0.76 μM for TREK1. GI-530159 displays selectivity for TREK1/2 over TRAAK, TASK3 and other potassium channels. GI-530159 reduces rat dorsal root ganglion neuron excitability<sup>[1]</sup>.

### REFERENCES

[1]. Loucif AJC, et al. GI-530159, a novel, selective, mechanosensitive two-pore-domain potassium (K<sub>2P</sub>) channel opener, reduces rat dorsal root ganglion neuron excitability. Br J Pharmacol. 2018;175(12):2272-2283.

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**Caution: Product has not been fully validated for medical applications. For research use only.**

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: [tech@MedChemExpress.com](mailto:tech@MedChemExpress.com)

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA