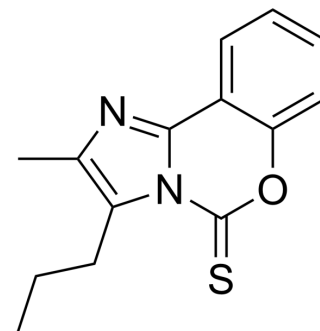


## NaV1.2/1.6 channel blocker-1

<b>Cat. No.:</b>	HY-152166
<b>CAS No.:</b>	1199944-04-1
<b>Molecular Formula:</b>	C <sub>14</sub> H <sub>14</sub> N <sub>2</sub> OS
<b>Molecular Weight:</b>	258.34
<b>Target:</b>	Sodium Channel
<b>Pathway:</b>	Membrane Transporter/Ion Channel
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	NaV1.2/1.6 channel blocker-1 is a potent NaV1.2/1.6 channel blocker, with IC <sub>50</sub> s of 9.8 and 24.4 μM for rNaV1.6 and hNaV1.2, respectively. NaV1.2/1.6 channel blocker-1 can be used for the research of generalized epilepsy <sup>[1]</sup> .	
<b>IC<sub>50</sub> &amp; Target</b>	rNaV1.6 9.8 μM (IC <sub>50</sub> )	hNaV1.2 24.4 μM (IC <sub>50</sub> )
<b>In Vitro</b>	NaV1.2/1.6 channel blocker-1 (compound 13) (1-100 μM) blocks the NaV1.6 and NaV1.2 sodium currents by 48.5% and 34.0% in HEK 293 stably expressing human NaV1.2 and rat NaV1.6 isoforms, respectively <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	

### REFERENCES

[1]. Rivara M, et, al. Biological Evaluation of Imidazobenzoxazines, Imidazobenzoxazin-5-ones and Imidazobenzoxazin-5-thiones as Sodium Channel Blockers. Letters in Drug Design & Discovery, 2014, 11, 90-97.

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

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