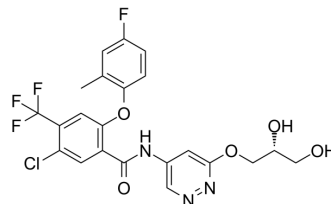


## Nav1.8-IN-7

Cat. No.:	HY-160588
CAS No.:	2761181-58-0
Molecular Formula:	C <sub>22</sub> H <sub>18</sub> ClF <sub>4</sub> N <sub>3</sub> O <sub>5</sub>
Molecular Weight:	515.84
Target:	Sodium Channel; Potassium Channel
Pathway:	Membrane Transporter/Ion Channel
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Nav1.8-IN-7 (Example 116) is a selective Nav1.8 inhibitor. Nav1.8-IN-7 shows an inhibition of >50% with 100 nM for Nav1.8. Nav1.8-IN-7 inhibits hERG with an IC <sub>50</sub> of 15.6 μM. Nav1.8-IN-7 has the potential for pain research <sup>[1]</sup> .								
<b>In Vitro</b>	Nav1.8-IN-9 (Example 116; 1 μM) shows inhibitions of 1.2%, 3.7%, <1%, <1%, 2.6%, <1% for Nav1.1, Nav1.3, Nav1.4, Nav1.5, Nav1.6, and Nav1.7, respectively <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.								
<b>In Vivo</b>	Pharmacokinetic Parameters of Nav1.8-IN-7 in male Sprague-Dawley rats <sup>[1]</sup> . <table border="1" style="margin-left: auto; margin-right: auto;"> <thead> <tr> <th></th> <th>PO (100 mg/kg)</th> </tr> </thead> <tbody> <tr> <td>C<sub>max</sub> (ng/mL)</td> <td>2566</td> </tr> <tr> <td>AUC<sub>0-24</sub> (ng/mL·h)</td> <td>42454</td> </tr> <tr> <td>t<sub>1/2</sub> (h)</td> <td>8.7</td> </tr> </tbody> </table> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>		PO (100 mg/kg)	C <sub>max</sub> (ng/mL)	2566	AUC <sub>0-24</sub> (ng/mL·h)	42454	t <sub>1/2</sub> (h)	8.7
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### REFERENCES

[1]. Junfeng Ren, et al. A benzylamine or benzyl alcohol derivative and its use. CN114031518A.

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

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