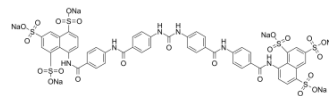


## NF279

Cat. No.:	HY-D0976
CAS No.:	202983-32-2
Molecular Formula:	C <sub>49</sub> H <sub>30</sub> N <sub>6</sub> Na <sub>6</sub> O <sub>23</sub> S <sub>6</sub>
Molecular Weight:	1401.12
Target:	P2X Receptor; HIV
Pathway:	Membrane Transporter/Ion Channel; Anti-infection
Storage:	Please store the product under the recommended conditions in the COA.



### BIOLOGICAL ACTIVITY

Description	NF279 is a potent selective and reversible <b>P2X1 receptor</b> antagonist, with an IC <sub>50</sub> of 19 nM. NF279 displays good selectivity over P2X2, P2X3 (IC <sub>50</sub> =1.62 μM), P2X4 (IC <sub>50</sub> >300 μM). NF279 is a dual HIV-1 coreceptor inhibitor that interferes with the functional engagement of CCR5 and CXCR4 by Env <sup>[1][2]</sup> .
IC <sub>50</sub> & Target	HIV

### REFERENCES

- [1]. Rettinger J, Schmalzing G, Damer S, Müller G, Nickel P, Lambrecht G. The suramin analogue NF279 is a novel and potent antagonist selective for the P2X(1) receptor. *Neuropharmacology*. 2000;39(11):2044-2053.
- [2]. Giroud C, et al. P2X1 Receptor Antagonists Inhibit HIV-1 Fusion by Blocking Virus-Coreceptor Interactions. *J Virol*. 2015;89(18):9368-9382.

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

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