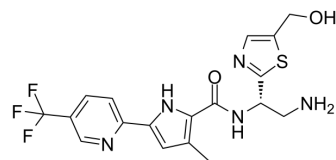


## NBD-14270

<b>Cat. No.:</b>	HY-139989
<b>CAS No.:</b>	2411819-82-2
<b>Molecular Formula:</b>	C <sub>18</sub> H <sub>18</sub> F <sub>3</sub> N <sub>5</sub> O <sub>2</sub> S
<b>Molecular Weight:</b>	425.43
<b>Target:</b>	HIV
<b>Pathway:</b>	Anti-infection
<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

<b>In Vitro</b>	DMSO : 100 mg/mL (235.06 mM; Need ultrasonic)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	<b>Preparing Stock Solutions</b>	1 mM	2.3506 mL	11.7528 mL	23.5056 mL
		5 mM	0.4701 mL	2.3506 mL	4.7011 mL
		10 mM	0.2351 mL	1.1753 mL	2.3506 mL
Please refer to the solubility information to select the appropriate solvent.					
<b>In Vivo</b>	<ol style="list-style-type: none"> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 40% PEG300 &gt;&gt; 5% Tween-80 &gt;&gt; 45% saline Solubility: ≥ 2.5 mg/mL (5.88 mM); Clear solution</li> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (5.88 mM); Clear solution</li> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 90% corn oil Solubility: ≥ 2.5 mg/mL (5.88 mM); Clear solution</li> </ol>				

### BIOLOGICAL ACTIVITY

<b>Description</b>	NBD-14270, a pyridine analogue, is a potent HIV-1 entry antagonist with an IC <sub>50</sub> of 180 nM against 50 HIV-1 Env-pseudotyped viruses. NBD-14270 binds to HIV-1 gp120 and shows potent antiviral activity. NBD-14270 shows low cytotoxicity (CC <sub>50</sub> >100 μM) <sup>[1][2]</sup> .
<b>IC<sub>50</sub> &amp; Target</b>	HIV-1
<b>In Vitro</b>	NBD-14270 has anti-HIV-1 activity (IC <sub>50</sub> =0.16 μM) and Cytotoxicity (CC <sub>50</sub> =109.3 μM) in single-cycle (TZM-bl Cells) assay <sup>[1]</sup> . NBD-14270 does not induce toxicity in the U87-CD4-CXCR4 cell line at the doses used for this assay <sup>[1]</sup> .

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MCE has not independently confirmed the accuracy of these methods. They are for reference only.

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## REFERENCES

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- [1]. Francesca Curreli, et al. Preclinical Optimization of gp120 Entry Antagonists as anti-HIV-1 Agents with Improved Cytotoxicity and ADME Properties through Rational Design, Synthesis, and Antiviral Evaluation. *J Med Chem.* 2020 Feb 27;63(4):1724-1749.
- [2]. Natalie Losada, et al. HIV-1 gp120 Antagonists Also Inhibit HIV-1 Reverse Transcriptase by Bridging the NNRTI and NRTI Sites. *J Med Chem.* 2021 Nov 25;64(22):16530-16540.
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**Caution: Product has not been fully validated for medical applications. For research use only.**

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