

产品名称: PFI-2 (hydrochloride)
产品别名: (R)-PFI-2 hydrochloride

生物活性:

Description	PFI-2 hydrochloride is a first-in-class, potent, highly selective, and cell-active inhibitor of the methyltransferase activity of SETD7 with IC50 of 2 nM, 500 fold active than (S)-PFI-2. IC50 value: 2 nM [1] Target: SETD7 (R)-PFI-2 is highly selective (>1,000-fold) for SETD7, over a panel of 18 other human protein methyltransferases and DNMT1, and was shown to be inactive against 134 additional ion channel, GPCR, and enzyme targets (<35% inhibition at 10 μ M). (R)-PFI-2 binds to SETD7 only in the presence of SAM. PFI-766, a biotinylated variant of (R)-PFI-2 that retains the ability to bind and inhibit SETD7 (IC50 110 \pm 26 nM in our in vitro enzymatic assay). PFI-766 engagement of endogenous SETD7 was also confirmed by mass spectrometry that supported the high specificity of the compound for endogenous SETD7.																					
In Vitro: DMSO : \geq 32 mg/mL (59.70 mM) H ₂ O : 5.56 mg/mL (10.37 mM; Need ultrasonic) * " \geq " means soluble, but saturation unknown. Preparing Stock Solutions	<table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th rowspan="2"></th> <th>Solvent Concentration</th> <th>Mass</th> <th>1 mg</th> <th>5 mg</th> <th>10 mg</th> </tr> </thead> <tbody> <tr> <td>1 mM</td> <td></td> <td>1.8657 mL</td> <td>9.3287 mL</td> <td>18.6574 mL</td> </tr> <tr> <td>5 mM</td> <td></td> <td>0.3731 mL</td> <td>1.8657 mL</td> <td>3.7315 mL</td> </tr> <tr> <td>10 mM</td> <td></td> <td>0.1866 mL</td> <td>0.9329 mL</td> <td>1.8657 mL</td> </tr> </tbody> </table> <p>*请根据产品在不同溶剂中的溶解度选择合适的溶剂配制储备液。一旦配成溶液，请分装保存，避免反复冻融造成的产品失效。</p> <p>储备液的保存方式和期限 -80°C, 6 months; -20°C, 1 month。 -80°C 储存时，请在 6 个月内使用，-20°C 储存时，请在 1 个月内使用。</p> <p>In Vivo:</p> <p>请根据您的实验动物和给药方式选择适当的溶解方案。以下溶解方案都请先按照 In Vitro 方式配制澄清的储备液，再依次添加助溶剂：</p> <p>——为保证实验结果的可靠性，澄清的储备液可以根据储存条件，适当保存；体内实验的工作液，建议您现用现配，当天使用；以下溶剂前显示的百分比是指该溶剂在您配制终溶液中的体积占比；如在配制过程中出现沉淀、析出现象，可以通过加热和/或超声的方式助溶。</p> <p>1. 请依序添加每种溶剂： 10% DMSO → 40% PEG300 → 5% Tween-80 → 45% saline Solubility: \geq 2.5 mg/mL (4.66 mM); Clear solution 此方案可获得 \geq 2.5 mg/mL (4.66 mM, 饱和度未知) 的澄清溶液。 以 1 mL 工作液为例，取 100 μL 25.0 mg/mL 的澄清 DMSO 储备液加到 400 μL PEG300 中，混合均匀；向上述体系中加入 50 μL Tween-80，混合均匀；然后继续加入 450 μL 生理盐水定容至 1 mL。</p> <p>2. 请依序添加每种溶剂： 10% DMSO → 90% (20% SBE-β-CD in saline) Solubility: \geq 2.5 mg/mL (4.66 mM); Clear solution 此方案可获得 \geq 2.5 mg/mL (4.66 mM, 饱和度未知) 的澄清溶液。 以 1 mL 工作液为例，取 100 μL 25.0 mg/mL 的澄清 DMSO 储备液加到 900 μL 20% 的 SBE-β-CD 生理盐水溶液中，混合均匀。</p>		Solvent Concentration	Mass	1 mg	5 mg	10 mg	1 mM		1.8657 mL	9.3287 mL	18.6574 mL	5 mM		0.3731 mL	1.8657 mL	3.7315 mL	10 mM		0.1866 mL	0.9329 mL	1.8657 mL
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Solvent&Solubility																						

	<p>3.请依序添加每种溶剂: 10% DMSO →90% corn oil Solubility: ≥ 2.5 mg/mL (4.66 mM); Clear solution</p> <p>此方案可获得 ≥ 2.5 mg/mL (4.66 mM, 饱和度未知) 的澄清溶液, 此方案不适用于实验周期在半个月以上的实验。</p> <p>以 1 mL 工作液为例, 取 100 μL 25.0 mg/mL 的澄清 DMSO 储备液加到 900 μL 玉米油中, 混合均匀。</p>
References	[1]. Barsyte-Lovejoy D, et al. (R)-PFI-2 is a potent and selective inhibitor of SETD7 methyltransferase activity in cells. Proc Natl Acad Sci U S A. 2014 Sep 2;111(35):12853-8.



源叶生物