

产品名称: **HPOB**

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**生物活性:**

<b>Description</b>	HPOB is a highly potent and selective inhibitor of histone deacetylase 6 (HDAC6) with IC <sub>50</sub> of 56 nM, >30 fold less potent against other HDACs. target: HDAC6 [1] IC <sub>50</sub> : 56nM [1] In vitro: HPOB causes growth inhibition of normal and transformed cells but does not induce cell death. HPOB enhances the effectiveness of DNA-damaging anticancer drugs in transformed cells but not normal cells. [1] Neuroprotective effect of HPOB demonstrated the crucial role of HDAC6 inhibition in reducing Cort-induced apoptosis in PC12 cells. Pre-treatment with HPOB remarkably reduces Cort-induced cytotoxicity and confirms the anti-apoptotic effect of HPOB via the caspase-3 activity assay and H33342/PI and TUNEL double staining. [2] In vivo: on corticosterone (Cort)-induced apoptosis and explores the possible mechanism of action of HPOB in rat adrenal pheochromocytoma (PC12) cells, which possesses typical neuron features and expresses high levels of glucocorticoid receptors. [2]				
<b>IC<sub>50</sub> &amp; Target</b>	HDAC6	HDAC3/NCOR2	HDAC8	HDAC1	HDAC10
	0.056 μM (IC <sub>50</sub> )	1.7 μM (IC <sub>50</sub> )	2.8 μM (IC <sub>50</sub> )	2.9 μM (IC <sub>50</sub> )	3.0 μM (IC <sub>50</sub> )
	HDAC2				
	4.4 μM (IC <sub>50</sub> )				
<b>Solvent&amp;Solubility</b>	<b>In Vitro:</b> <b>DMSO : 50 mg/mL (159.06 mM; Need ultrasonic)</b>				
	<b>Preparing Stock Solutions</b>	<div>Solvent / Mass / Concentration</div>	<b>1 mg</b>	<b>5 mg</b>	<b>10 mg</b>
		1 mM	3.1813 mL	15.9063 mL	31.8127 mL
		5 mM	0.6363 mL	3.1813 mL	6.3625 mL
		10 mM	0.3181 mL	1.5906 mL	3.1813 mL
	*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。				
	储备液的保存方式和期限 -80℃, 6 months; -20℃, 1 month. -80℃ 储存时，请在 6 个月内使用，-20℃ 储存时，请在 1 个月内使用。				
	<b>In Vivo:</b>				
	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 <b>In Vitro</b> 方式配制澄清的储备液，再依次添加助溶剂：				
	——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				
	<div>1.请依序添加每种溶剂： 10% DMSO→40% PEG300 →5% Tween-80 → 45% saline Solubility: ≥ 2.5 mg/mL (7.95 mM); Clear solution 此方案可获得 ≥ 2.5 mg/mL (7.95 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。</div>				
	<div>2.请依序添加每种溶剂： 10% DMSO→ 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (7.95 mM); Clear solution</div>				

	<p>此方案可获得 <math>\geq 2.5</math> mg/mL (7.95 mM, 饱和度未知) 的澄清溶液。</p> <p>以 1 mL 工作液为例, 取 100 <math>\mu</math>L 25.0 mg/mL 的澄清 DMSO 储备液加到 900 <math>\mu</math>L 20% 的 SBE-<math>\beta</math>-CD 生理盐水水溶液中, 混合均匀。</p> <p>3.请依序添加每种溶剂: 10% DMSO <math>\rightarrow</math> 90% corn oil</p> <p>Solubility: <math>\geq 2.5</math> mg/mL (7.95 mM); Clear solution</p> <p>此方案可获得 <math>\geq 2.5</math> mg/mL (7.95 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 <math>\mu</math>L 25.0 mg/mL 的澄清 DMSO 储备液加到 900 <math>\mu</math>L 玉米油中, 混合均匀。</p>
References	<p>[1]. <a href="#">Lee JH et al. Development of a histone deacetylase 6 inhibitor and its biological effects. Proc Natl Acad Sci U S A. 2013 Sep 24;110(39):15704-9.</a></p> <p>[2]. <a href="#">Li ZY et al. HPOB, an HDAC6 inhibitor, attenuates corticosterone-induced injury in rat adrenal pheochromocytoma PC12 cells by inhibiting mitochondrial GR translocation and the intrinsic apoptosis pathway. Neurochem Int. 2016 Oct;99:239-51.</a></p>

源叶生物