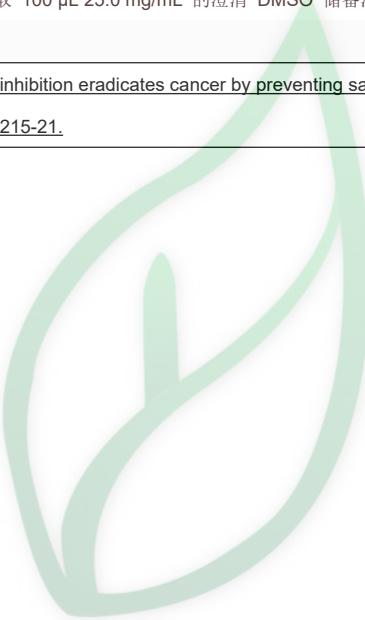


产品名称: TH588

产品别名: TH588

生物活性:																							
Description	TH588 is first-in-class nudix hydrolase family inhibitor that potently and selectively engage and inhibit the MTH1 (IC_{50} = 5 nM).																						
IC₅₀ & Target	IC50: 5 nM (MTH1)[1]																						
In Vitro	TH588 (2-10μM; 7-10 days) selectively and effectively kills U2OS, HeLa, MDA-MB-231, MCF-7, SW480, and SW620 cells with IC50s of 1.38, 0.83, 1.03, 1.08, 1.72, 0.8 μM[1].																						
	Cell Viability Assay[1]																						
	Cell Line: U2OS, HeLa, MDA-MB-231, MCF-7, SW480, SW620, VH10, HDFn cells																						
	Concentration: 2, 4, 6, 8, 10 μM																						
	Incubation Time: 7-10 days																						
In Vivo	Result: Selectively and effectively killed U2OS, HeLa, MDA-MB-231, MCF-7, SW480, and SW620 cells with IC ₅₀ s of 1.38, 0.83, 1.03, 1.08, 1.72, 0.8 μM, but was less toxic to several primary or immortalized cells.																						
	TH588 (30 mg/kg; s.c.; once daily for 35 days) reduces tumour growth in SW480 xenograft cancer model[1].																						
	Animal Model: 5-6 weeks female SCID mice (SW480 xenograft cancer model)[1]																						
	Dosage: 30 mg/kg																						
	Administration: Subcutaneous injection; once daily for 35 days																						
Solvent&Solubility	Result: Reduced tumour growth in SW480 xenograft cancer model.																						
	In Vitro:																						
	DMSO : 16 mg/mL (54.21 mM; Need ultrasonic and warming)																						
	<table border="1"><thead><tr><th rowspan="2">Preparing Stock Solutions</th><th>Solvent</th><th>Mass</th><th rowspan="2">1 mg</th><th rowspan="2">5 mg</th><th rowspan="2">10 mg</th></tr><tr><th>Concentration</th><th></th></tr></thead><tbody><tr><td></td><td>1 mM</td><td>3.3879 mL</td><td>16.9394 mL</td><td>33.8788 mL</td></tr><tr><td></td><td>5 mM</td><td>0.6776 mL</td><td>3.3879 mL</td><td>6.7758 mL</td></tr><tr><td></td><td>10 mM</td><td>0.3388 mL</td><td>1.6939 mL</td><td>3.3879 mL</td></tr></tbody></table>	Preparing Stock Solutions	Solvent	Mass	1 mg	5 mg	10 mg	Concentration			1 mM	3.3879 mL	16.9394 mL	33.8788 mL		5 mM	0.6776 mL	3.3879 mL	6.7758 mL		10 mM	0.3388 mL	1.6939 mL
Preparing Stock Solutions	Solvent		Mass	1 mg				5 mg	10 mg														
	Concentration																						
	1 mM	3.3879 mL	16.9394 mL	33.8788 mL																			
	5 mM	0.6776 mL	3.3879 mL	6.7758 mL																			
	10 mM	0.3388 mL	1.6939 mL	3.3879 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°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。																						
	In Vivo:																						
	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:																						
	——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																						
	1.请依序添加每种溶剂: 10% DMSO→40% PEG300 →5% Tween-80 → 45% saline Solubility: ≥ 2.5 mg/mL (8.47 mM); Clear solution 此方案可获得 ≥ 2.5 mg/mL (8.47 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																						

	<p>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: $\geq 2.5 \text{ mg/mL}$ (8.47 mM); Clear solution 此方案可获得 $\geq 2.5 \text{ mg/mL}$ (8.47 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> <p>3.请依序添加每种溶剂： 10% DMSO → 90% corn oil Solubility: $\geq 2.5 \text{ mg/mL}$ (8.47 mM); Clear solution 此方案可获得 $\geq 2.5 \text{ mg/mL}$ (8.47 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。 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]. Gad H, et al. MTH1 inhibition eradicates cancer by preventing sanitation of the dNTP pool. Nature. 2014 Apr 10;508(7495):215-21.



源叶生物