



上海源叶生物科技有限公司
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产品名称: CAY10650

产品别名: CAY10650

生物活性:

Description	CAY10650 is a highly potent cytosolic phospholipase A2 α (cPLA2 α) inhibitor with an IC50 value of 12 nM. IC50 value: 12 nM Target: cPLA2 CAY10650 is a highly potent (IC50 = 12 nM) cPLA2 α inhibitor. It demonstrates strong anti-inflammatory effects when applied topically at a dose of 0.1 mg/ear in a mouse model of acute irritant contact dermatitis. Chinese hamsters ($n = 6/\text{group}$) were infected with parasite-laden contact lenses and treated with cPLA2 α inhibitors (AACOCF3 and CAY10650) 50 $\mu\text{g}/5 \mu\text{l}$ was injected with topical eye-drop under the contact lens of an infected cornea three times a day for 6 days and topically on days 7–14 postinfection. Animals were anesthetized and sacrificed 15 days after application of cPLA2 α inhibitors. Treatment with the AACOCF3 and CAY10650 had a profound effect on the severity and chronicity of keratitis. In addition, hamsters treated with AACOCF3 had significantly less severe keratitis as compared with CAY10650 group.																									
In Vitro: DMSO : $\geq 43 \text{ mg/mL}$ (91.20 mM) * " \geq " means soluble, but saturation unknown.	<table border="1"><thead><tr><th rowspan="2">Preparing Stock Solutions</th><th>Solvent</th><th>Mass</th><th>Concentration</th><th>1 mg</th><th>5 mg</th><th>10 mg</th></tr></thead><tbody><tr><td>1 mM</td><td>2.1209 mL</td><td></td><td>10.6045 mL</td><td>21.2089 mL</td><td></td></tr><tr><td>5 mM</td><td>0.4242 mL</td><td></td><td>2.1209 mL</td><td>4.2418 mL</td><td></td></tr><tr><td>10 mM</td><td>0.2121 mL</td><td></td><td>1.0604 mL</td><td>2.1209 mL</td><td></td></tr></tbody></table>	Preparing Stock Solutions	Solvent	Mass	Concentration	1 mg	5 mg	10 mg	1 mM	2.1209 mL		10.6045 mL	21.2089 mL		5 mM	0.4242 mL		2.1209 mL	4.2418 mL		10 mM	0.2121 mL		1.0604 mL	2.1209 mL	
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Solvent&Solubility	*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: $\geq 2.5 \text{ mg/mL}$ (5.30 mM); Clear solution 此方案可获得 $\geq 2.5 \text{ mg/mL}$ (5.30 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。 2.请依序添加每种溶剂: 10% DMSO → 90% (20% SBE-β-CD in saline) Solubility: $\geq 2.5 \text{ mg/mL}$ (5.30 mM); Clear solution 此方案可获得 $\geq 2.5 \text{ mg/mL}$ (5.30 mM, 饱和度未知) 的澄清溶液。																									



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	<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 20% 的 SBE-β-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 → 90% corn oil Solubility: ≥ 2.5 mg/mL (5.30 mM); Clear solution 此方案可获得 ≥ 2.5 mg/mL (5.30 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]. Tripathi T, et al. Role of phospholipase A 2(PLA2) inhibitors in attenuating apoptosis of the corneal epithelial cells and mitigation of Acanthamoeba keratitis. <i>Exp Eye Res.</i> 2013 Aug;113:182-91.



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