

产品名称: HJC0350

产品别名: HJC 0350

生物活性:																									
Description	HJC0350 is a potent and specific EPAC2 antagonist with an IC ₅₀ of 0.3 μM.																								
IC₅₀ & Target	IC50: 0.3 μM (EPAC2)[1]																								
In Vitro	HJC0350 has an apparent IC50 value of 0.3 μM for competing with 8-NBD-cAMP binding of EPAC2, and is about 133-fold more potent than cAMP. HJC0350 is found not to inhibit EPAC1-mediated Rap1-GDP exchange activity at 25 μM in the presence of equal concentration of cAMP, indicating that it is EPAC2-specific antagonists. Pretreatment of HEK293/EPAC2-FL cells with 10 μM HJC0350 fully blocks the 007-AM induced decrease of FRET[1].																								
In Vitro: DMSO : 33.33 mg/mL (120.16 mM; Need ultrasonic)	<table border="1"><thead><tr><th rowspan="2">Preparin Stock Solutions</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>3.6052 mL</td><td>18.0258 mL</td><td>36.0516 mL</td></tr><tr><td>5 mM</td><td></td><td>0.7210 mL</td><td>3.6052 mL</td><td>7.2103 mL</td></tr><tr><td>10 mM</td><td></td><td>0.3605 mL</td><td>1.8026 mL</td><td>3.6052 mL</td></tr></tbody></table>				Preparin Stock Solutions	Solvent Concentration	Mass	1 mg	5 mg	10 mg	1 mM		3.6052 mL	18.0258 mL	36.0516 mL	5 mM		0.7210 mL	3.6052 mL	7.2103 mL	10 mM		0.3605 mL	1.8026 mL	3.6052 mL
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Solvent&Solubility	<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 → 90% corn oil</p> <p>Solubility: ≥ 2.5 mg/mL (9.01 mM); Clear solution</p> <p>此方案可获得 ≥ 2.5 mg/mL (9.01 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]. Chen H, et al. Identification and characterization of small molecules as potent and specific EPAC2 antagonists. J Med Chem. 2013 Feb 14;56(3):952-62.																								