



上海源叶生物科技有限公司
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产品名称: CID 16020046

产品别名: CID 16020046

生物活性:

Description	<p>CID 16020046 is a potent and selective GPR55(LPI receptor) antagonist; inhibits GPR55 constitutive activity with IC50 of 0.15 uM. IC50 value: 0.15 uM [1] Target: GPR55 antagonist In yeast cells expressing human GPR55, CID16020046 antagonized agonist-induced receptor activation. In human embryonic kidney(HEK293) cells stably expressing human GPR55, the compound behaved as an antagonist on LPI-mediated Ca2+ release and extracellular signal-regulated kinases activation, but not in HEK293 cells expressing cannabinoid receptor 1 or 2. CID16020046 concentration dependently inhibited LPI-induced activation of nuclear factor of activated T-cells (NFAT), nuclear factor k of activated B cells (NF-κB) and serum response element, translocation of NFAT and NF-κB, and GPR55 internalization. It reduced LPI-induced wound healing in primary human lung microvascular endothelial cells and reversed LPI-inhibited platelet aggregation.</p>																												
In Vitro: DMSO : 100 mg/mL (235.05 mM; Need ultrasonic)	<table border="1"><thead><tr><th rowspan="2">Preparing Stock Solutions</th><th>Solvent Mass</th><th>1 mg</th><th>5 mg</th><th>10 mg</th></tr><tr><th>Concentration</th><th></th><th></th><th></th></tr></thead><tbody><tr><td>1 mM</td><td>2.3505 mL</td><td>11.7525 mL</td><td>23.5051 mL</td><td></td></tr><tr><td>5 mM</td><td>0.4701 mL</td><td>2.3505 mL</td><td>4.7010 mL</td><td></td></tr><tr><td>10 mM</td><td>0.2351 mL</td><td>1.1753 mL</td><td>2.3505 mL</td><td></td></tr></tbody></table>					Preparing Stock Solutions	Solvent Mass	1 mg	5 mg	10 mg	Concentration				1 mM	2.3505 mL	11.7525 mL	23.5051 mL		5 mM	0.4701 mL	2.3505 mL	4.7010 mL		10 mM	0.2351 mL	1.1753 mL	2.3505 mL	
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<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>	In Vivo:	<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>																									
Solvent&Solubility	<p>1.请依序添加每种溶剂: 10% DMSO→40% PEG300 →5% Tween-80 → 45% saline</p>	<p>Solubility: ≥ 2.5 mg/mL (5.88 mM); Clear solution</p>	<p>此方案可获得 ≥ 2.5 mg/mL (5.88 mM, 饱和度未知) 的澄清溶液。</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 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)</p>	<p>Solubility: ≥ 2.5 mg/mL (5.88 mM); Clear solution</p>	<p>此方案可获得 ≥ 2.5 mg/mL (5.88 mM, 饱和度未知) 的澄清溶液。</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 20% 的 SBE-β-CD 生理</p>																									



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	<p>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</p> <p>Solubility: ≥ 2.5 mg/mL (5.88 mM); Clear solution</p> <p>此方案可获得 ≥ 2.5 mg/mL (5.88 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]. Kargl J, et al. A selective antagonist reveals a potential role of G protein-coupled receptor 55 in platelet and endothelial cell function. <i>J Pharmacol Exp Ther.</i> 2013 Jul;346(1):54-66.



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