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产品名称: **JZL195**

产品别名: **JZL195**

**生物活性:**

<b>Description</b>	JZL195 is a selective and efficacious dual fatty acid amide hydrolase (FAAH) and monoacylglycerol lipase (MAGL) inhibitor with IC50s of 2 and 4 nM, respectively[1].																												
<b>IC<sub>50</sub> &amp; Target</b>	IC50: 2 nM (FAAH), 4 nM (MAGL)[1]																												
<b>In Vitro</b>	JZL195 produces near-complete blockade of FP-Rh labeling of both mouse brain FAAH and MAGL at concentrations as low as 100 nM (IC50 values of 13 and 19 nM, respectively)[1]. JZL195 inhibits rat and human FAAH and MAGL enzymes with IC50 values in the range of ≈10-100 nM based on competitive ABPP assays[1].																												
<b>In Vivo</b>	JZL195 (20 mg/kg; i.p.) produces an antinociceptive response in the tail immersion assay[1]. <b>Animal Model:</b> Male C57BL/6J mice[1] <b>Dosage:</b> 20 mg/kg <b>Administration:</b> Intraperitoneal injection <b>Result:</b> Produced a much greater antinociceptive response in the tail immersion assay compared with inhibitors of either FAAH or MAGL alone.																												
<b>Solvent&amp;Solubility</b>	<p><b>In Vitro:</b> DMSO : 50 mg/mL (115.35 mM; Need ultrasonic)</p> <table border="1"><thead><tr><th rowspan="2">Preparing Stock Solutions</th><th>Solvent</th><th>Mass</th><th>Concentration</th><th></th></tr><tr><th></th><th>1 mg</th><th></th><th>5 mg</th><th>10 mg</th></tr></thead><tbody><tr><th>1 mM</th><td>2.3070 mL</td><td></td><td>11.5351 mL</td><td>23.0702 mL</td></tr><tr><th>5 mM</th><td>0.4614 mL</td><td></td><td>2.3070 mL</td><td>4.6140 mL</td></tr><tr><th>10 mM</th><td>0.2307 mL</td><td></td><td>1.1535 mL</td><td>2.3070 mL</td></tr></tbody></table> <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><b>In Vivo:</b></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 → 40% PEG300 → 5% Tween-80 → 45% saline</p> <p>Solubility: ≥ 2.5 mg/mL (5.77 mM); Clear solution</p> <p>此方案可获得 ≥ 2.5 mg/mL (5.77 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% corn oil</p>				Preparing Stock Solutions	Solvent	Mass	Concentration			1 mg		5 mg	10 mg	1 mM	2.3070 mL		11.5351 mL	23.0702 mL	5 mM	0.4614 mL		2.3070 mL	4.6140 mL	10 mM	0.2307 mL		1.1535 mL	2.3070 mL
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	<p>Solubility: <math>\geq 2.5 \text{ mg/mL}</math> (5.77 mM); Clear solution</p> <p>此方案可获得 <math>\geq 2.5 \text{ mg/mL}</math> (5.77 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\text{L}</math> 25.0 mg/mL 的澄清 DMSO 储备液加到 900 <math>\mu\text{L}</math> 玉米油中, 混合均匀。</p>
<b>References</b>	<p>[1]. Long JZ, et al. Dual blockade of FAAH and MAGL identifies behavioral processes regulated by endocannabinoid crosstalk in vivo. Proc Natl Acad Sci U S A. 2009 Dec 1;106(48):20270-5.</p> <p>[2]. Anderson WB, et al. Actions of the dual FAAH/MAGL inhibitor JZL195 in a murine inflammatory pain model. Neuropharmacology. 2014 Jun;81:224-30.</p>



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