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产品名称: **AZD2098**  
 产品别名: **AZD2098**

生物活性:																									
<b>Description</b>	AZD2098 is a potent and selective CC-chemokine receptor 4 (CCR4) inhibitor with pIC <sub>50</sub> s of 7.8, 8.0, 8.0 and 7.6 for human, rat, mouse and dog respectively, used for asthma research[1][2].																								
<b>IC<sub>50</sub> &amp; Target</b>	CCR4 [1]																								
<b>In Vitro</b>	AZD2098 potently inhibits chemokine-induced cellular responses, with pIC <sub>50</sub> of 7.5 and 6.3 against CCL22-induced Ca <sup>2+</sup> influx in hCCR4-expressing CHO cells and CCL17- or CCL22-induced chemotaxis of primary human Th2 cells respectively[1].																								
<b>In Vivo</b>	AZD2098 (73.5-5.0 µg/kg; p.o.; BID; twice a day; 1 hour before and every 12 hours after antigen challenge) exhibits efficacy against antigen-induced inflammatory response among ovalbumin-sensitized rats, and the changes are first visible at a dose of 0.22 µmol/kg and maximal at 7.5 µmol/kg[1].																								
	<b>Animal Model:</b> Brown-Norway rats[1]																								
	<b>Dosage:</b> 73.5 µg/kg, 250.6 µg/kg, 735.2 µg/kg, 1.0 mg/kg, 2.5 mg/kg, 5.0 mg/kg																								
	<b>Administration:</b> Oral administration; twice a day; 1 hour before and every 12 hours after antigen challenge																								
<b>Result:</b>	Exhibits efficacy against antigen-induced inflammatory response among ovalbumin-sensitized rats.																								
<b>Solvent&amp;Solubility</b>	<b>In Vitro:</b> <b>DMSO : 100 mg/mL (299.24 mM; Need ultrasonic)</b> <b>H<sub>2</sub>O : &lt; 0.1 mg/mL (insoluble)</b>																								
	<table border="1"> <thead> <tr> <th rowspan="2">Preparing</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 rowspan="3"><b>Stock Solutions</b></td> <td>1 mM</td> <td></td> <td>2.9924 mL</td> <td>14.9620 mL</td> <td>29.9240 mL</td> </tr> <tr> <td>5 mM</td> <td></td> <td>0.5985 mL</td> <td>2.9924 mL</td> <td>5.9848 mL</td> </tr> <tr> <td>10 mM</td> <td></td> <td>0.2992 mL</td> <td>1.4962 mL</td> <td>2.9924 mL</td> </tr> </tbody> </table>	Preparing	Solvent	Mass	1 mg	5 mg	10 mg	Concentration		<b>Stock Solutions</b>	1 mM		2.9924 mL	14.9620 mL	29.9240 mL	5 mM		0.5985 mL	2.9924 mL	5.9848 mL	10 mM		0.2992 mL	1.4962 mL	2.9924 mL
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*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。																									
<b>In Vivo:</b> 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 (7.48 mM); Clear solution 此方案可获得 ≥ 2.5 mg/mL (7.48 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																									



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	<p>向上述体系中加入 50 <math>\mu</math>L Tween-80, 混合均匀; 然后继续加入 450 <math>\mu</math>L 生理盐水定容至 1 mL。</p> <p>2.请依序添加每种溶剂: 10% DMSO <math>\rightarrow</math>90% corn oil</p> <p>Solubility: <math>\geq</math> 2.5 mg/mL (7.48 mM); Clear solution</p> <p>此方案可获得 <math>\geq</math> 2.5 mg/mL (7.48 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</math>L 25.0 mg/mL 的澄清 DMSO 储备液加到 900 <math>\mu</math>L 玉米油中, 混合均匀。</p>
<b>References</b>	<p>[1]. Kindon N, et al. Discovery of AZD-2098 and AZD-1678, Two Potent and Bioavailable CCR4 Receptor Antagonists. ACS Med Chem Lett. 2017 Sep 1;8(9):981-986.</p> <p>[2]. Asher Mullard. Cancer charity sees success re-prioritizing industry's shelved compounds. Nat Rev Drug Discov. 2014 May;13(5):319-21.</p>



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