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产品名称: AZD1283

产品别名: AZD1283

**生物活性:**

<b>Description</b>	AZD1283 is a potent antagonist of the P2Y12 receptor with EC50 of 3.0 ug/kg/min, TI >10; with binding IC50 of 11 nM. IC50 value: 3.0 ug/kg/min(EC50) [1] Target: P2Y12 receptor inhibitor AZD1283 dose-dependently induced increases in blood flow and inhibition of ADP-induced platelet aggregation with antithrombotic ED50 values of 3.0 and 10 µg/kg/min, respectively. The doses that induced a larger than 3-fold increase in bleeding time were 33 and 100 µg/kg/min for 3 and 13, respectively. Thus, the therapeutic index (TI) was ≥ 10 for both compounds. On the basis of these data, compound 3 was progressed into human clinical trials as candidate drug AZD1283.																					
<b>In Vitro:</b>  DMSO : 100 mg/mL (212.52 mM; Need ultrasonic)  H <sub>2</sub> O : < 0.1 mg/mL (insoluble)	<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>2.1252 mL</td><td>10.6261 mL</td><td>21.2522 mL</td></tr><tr><td>5 mM</td><td>0.4250 mL</td><td>2.1252 mL</td><td>4.2504 mL</td></tr><tr><td>10 mM</td><td>0.2125 mL</td><td>1.0626 mL</td><td>2.1252 mL</td></tr></tbody></table>	Preparing Stock Solutions	Solvent / Mass	1 mg	5 mg	10 mg	Concentration				1 mM	2.1252 mL	10.6261 mL	21.2522 mL	5 mM	0.4250 mL	2.1252 mL	4.2504 mL	10 mM	0.2125 mL	1.0626 mL	2.1252 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 (5.31 mM); Clear solution  此方案可获得 ≥ 2.5 mg/mL (5.31 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% corn oil  Solubility: ≥ 2.5 mg/mL (5.31 mM); Clear solution  此方案可获得 ≥ 2.5 mg/mL (5.31 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。																						



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**References**

- [1]. Bach P, et al. Lead optimization of ethyl 6-aminonicotinate acyl sulfonamides as antagonists of the P2Y12 receptor. separation of the antithrombotic effect and bleeding for candidate drug AZD1283. *J Med Chem.* 2013 Sep 12;56(17):7015-24.



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