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产品名称: **SD-06**
产品别名: **SD 0006**

生物活性:					
Description	SD 0006 (SD-06) is an orally active, selective, ATP-competitive and potent diaryl pyrazole inhibitor of p38α MAP kinase, with an IC50 of 110 nM for p38α[1][2].				
IC50 & Target	IC50: 110 nM (p38 MAPK)[1].				
In Vitro	SD 0006 clearly inhibits p38α as shown by the dose-dependent inhibition of phosphorylation of its endogenous Hsp27 substrate[1].				
In Vivo	SD 0006 (0-30 mg/kg) may be an effective alternative to steroids and biologics for RA therapy[1]. SD0006 (3.75, 7.5 and 15 mg/kg; p.o.; b.i.d.) is highly effective in attenuating SCW-induced inflammation as shown by the dose-dependent inhibition of paw swelling[1].				
	Animal Model:	8- to 12-week-old DBA/1 mice[1].			
	Dosage:	3.75, 7.5 and 15 mg/kg.			
	Administration:	Orally twice daily.			
	Result:	Inhibited the transcription of several inflammatory mediators to prevent joint swelling and bone destruction and to preserve bone density.			
Solvent&Solubility	In Vitro: DMSO : ≥ 29 mg/mL (72.89 mM) * "≥" means soluble, but saturation unknown.				
	Preparing Stock Solutions	<div><div>Solvent</div><div>Mass</div><div>Concentration</div></div>	1 mg	5 mg	10 mg
		1 mM	2.5134 mL	12.5672 mL	25.1345 mL
		5 mM	0.5027 mL	2.5134 mL	5.0269 mL
		10 mM	0.2513 mL	1.2567 mL	2.5134 mL
	*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℃，6 months；-20℃，1 month。 -80℃ 储存时，请在 6 个月内使用，-20℃ 储存时，请在 1 个月内使用。				
References	[1]. Burnette BL, et al. SD0006: a potent, selective and orally available inhibitor of p38 kinase. Pharmacology. 2009;84(1):42-60. [2]. Walker JK, et al. Identification of SD-0006, a potent diaryl pyrazole inhibitor of p38 MAP kinase. Bioorg Med Chem Lett. 2010 Apr 15;20(8):2634-8.				