

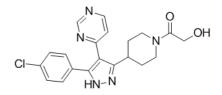
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## **Data Sheet**

Global Supplier of Chemical Probes, Inhibitors & Agonists.

Product Name	: SD0006
Cat. No.	: PC-42805
CAS No.	: 271576-80-8
Molecular Formula	: C <sub>20</sub> H <sub>20</sub> CIN <sub>5</sub> O <sub>2</sub>
Molecular Weight	: 397.8581
Target	: p38 MAPK
Solubility	: DMSO: ≥ 29 mg/mL
1. Burnette BL, et al. <b>Pharmacology</b> . 2009;84(1):42-60.	

2. Walker JK, et al. Bioorg Med Chem Lett. 2010 Apr 15;20(8):2634-8.



## **Biological Activity**

SD0006 (SD-0006, SD-06) is a potent, selective, ATP-competitive and orally bioavailable **p38** $\alpha$  inhibitor with Ki of 61 nM. SD0006 displays 8-fold selectivity over p38 $\beta$  (Ki=510 nM), and greater than 100-fold selectivity based upon IC50 ratios over the other MAPKs p38 $\gamma$ , p38 $\delta$ , ERK-2 and JNK-1, -2, and -3.

SD0006 potently inhibits tumor necrosis factor-alpha (TNFalpha) release, in vitro and in vivo, with IC50 of <200 nM. SD0006 is effective in the rat streptococcal-cell-wall-induced arthritis model, with dramatic protective effects on paw joint integrity and bone density as shown by radiographic analysis.

SD0006 also demonstrates good oral anti-inflammatory efficacy with excellent cross-species correlation between the rat, cynomolgus monkey, and human.

## References