Revision 1

e at -20C	SP600125		Cell Signaling
Store		Orders:	877-616-CELL (2355) orders@cellsignal.com
2	10 mg	Support:	877-678-TECH (8324)
17		Web:	info@cellsignal.com cellsignal.com
#81		3 Trask Lane Danvers M	assachusetts 01923 USA

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Background	Novel, potent, and selective JNK-1,-2, and -3 inhibitor, SP600125 is an ATP-competetive inhibitor effective on a range of kinases and enzymes. In cells, SP600125 caused a dose-dependent inhibition o the phosphorylation of c-Jun, the expression of inflammatory genes IL-2, COX-2, TNF-α, IFN-γ, and blocked the activation and differentiation of primary human CD4 cell cultures (1). SP600125 has also demonstrated inhibitory effects on tumor cell proliferation, endothelial cell migration, and tumor growth as well as blocking tumor and endothelial cells in the G ₂ phase of the cell cycle (2).
Molecular Formula	C ₁₄ H ₈ N ₂ O
Molecular Weight	220.23 g/mol
Purity	>99%
CAS	129-56-6
Solubility	Soluble in DMSO at 65mg/ml.
Storage	Store lyophilized or in solution at -20°C, desiccated. Protect from light. In lyophilized form, the chemica is stable for 24 months. Once in solution, use within 3 months to prevent loss of potency. Aliquot to avoid multiple freeze/thaw cycles.
Directions for Use	SP600125 is supplied as a lyophilized powder. For a 25 mM stock, reconstitute the 10 mg in 1.82 ml DMSO. Working concentrations and length of treatment can vary depending on the desired effect, but it is typically used as a pre-treatment at 25-50 μM for 15-45 minutes prior to treating with a stimulator. Soluble in DMSO at 65 mg/ml; poorly soluble in ethanol and water.
Background References	1. Bennett, B.L. et al. (2001) <i>Proc Natl Acad Sci U S A</i> 98, 13681-6. 2. Ennis, B.W. et al. (2005) <i>J Pharmacol Exp Ther</i> 313, 325-32.
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