MK2-IN-3

®

MedChemExpress

Cat. No.:	HY-131249
CAS No.:	724711-21-1
Molecular Formula:	$C_{21}H_{16}N_4O$
Molecular Weight:	340.38
Target:	MAPKAPK2 (MK2)
Pathway:	MAPK/ERK Pathway
Storage:	4°C, protect from light * In solvent : -80°C, 6 months; -20°C, 1 month (protect from light)

SOLVENT & SOLUBILITY

In Vitro	DMSO : 62.5 mg/mL (183.62 mM; Need ultrasonic)					
		Solvent Mass Concentration	1 mg	5 mg	10 mg	
	Preparing Stock Solutions	1 mM	2.9379 mL	14.6895 mL	29.3789 mL	
	Stock Solutions	5 mM	0.5876 mL	2.9379 mL	5.8758 mL	
		10 mM	0.2938 mL	1.4689 mL	2.9379 mL	
	Please refer to the solubility information to select the appropriate solvent.					

Description	MK2-IN-3 is a potent and selective inhibitor of MAPKAP-K2 (MK-2), with an IC ₅₀ of 8.5 nM. MK2-IN-3 shows selectivity for MK-2 over MK-3, MK-5, ERK2, MNK1, p38a (IC ₅₀ s=0.21, 0.081, 3.44, 5.7, and >100 μM, respectively) and MSK1, MSK2, CDK2, JNK2, IKK2 (IC ₅₀ s>200 μM). MK2-IN-3 can reduce TNFα production in both U937 cells and in vivo ^[1] .			
IC ₅₀ & Target	IC50: 8.5 nM (MK-2); 81 nM (MK-5); 210 nM (MK-3); 3.44 μ M (ERK2); 5.7 μ M (MNK1)^{[1]}			
In Vitro	MK2-IN-3 (compound 16) inhibits MK-2 and TNFα production in U937 cells, with IC ₅₀ s of 8.5 nM and 4.4 μM, respectively ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.			
In Vivo	MK2-IN-3 (compound 16) (20 mg/kg; a single p.o. before LPS challenge) inhibits 20% TNFα production in rat LPS (rLPS) model ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.			

CUSTOMER VALIDATION

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REFERENCES

[1]. Anderson DR, et, al. Pyrrolopyridine inhibitors of mitogen-activated protein kinase-activated protein kinase 2 (MK-2). J Med Chem. 2007 May 31;50(11):2647-54.

Caution: Product has not been fully validated for medical applications. For research use only.

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