H-89, Dihyd	lrochloride	С	Cell Signaling TECHNOLOGY®	
Stor		Orders:	877-616-CELL (2355) orders@cellsignal.com	
5.19 mg		Support:	877-678-TECH (8324)	
842		Web:	info@cellsignal.com cellsignal.com	
6		3 Trask Lane Danvers Mas	sachusetts 01923 USA	
For Research Use Only. No	ot for Use in Diagnostic Procedures.			
Background	H-89 is a potent selective inhibitor of cAMP dependent protein kinase (PKA). The <i>in vitro</i> IC50 of H-89 for PKA is approximately 50 nM and <i>in vivo</i> the inhibitiory effect on PKA substrate phosphorylation and related cellular functions range from 10 μ M to 30 μ M (1,2). In addition to PKA, H-89 also exhibits a moderate inhibitory effect on PKG and PKC μ , with IC50 in the 500 nM range (1,3). The inhibitory effect of H-89 is due to its competitive binding to the ATP pocket on the kinase catalytic subunit (4).			

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Molecular Formula	$C_{20}H_{20}BrN_3O_2S \cdot 2HCI$	
Molecular Weight	519.3 g/mol	
Purity	>99%	
CAS	127243-85-0	
Solubility	Soluble in DMSO at 100mg/ml and EtOH at 2mg/ml.	
Storage	Store lyophilized or in solution at -20°C, desiccated. Protect from light. In lyophilized form, the chemical 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	H-89 is supplied as 5.19 mg powder. Store at or below -20°C. Before usage, dissolve powder in 0.5 ml DMSO to make 20 mM H-89. For working concentrations of 10 μM-20 μM, dilute DMSO stock 1:2000 to 1:1000. Treat cells with the desired concentration for 30 minutes.	
Background References	1. Chijiwa, T. et al. (1990) <i>J Biol Chem</i> 265, 5267-72. 2. Meja, K.K. et al. (2004) <i>J Pharmacol Exp Ther</i> 309, 833-44. 3. Johannes, F.J. et al. (1995) <i>Eur J Biochem</i> 227, 303-7. 4. Engh, R.A. et al. (1996) <i>J Biol Chem</i> 271, 26157-64.	
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